



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 168245

TO: Jeffrey Fredman
Location: rem/2C89/2C18
Art Unit: 1637
Friday, October 14, 2005

Case Serial Number: 09/744097

From: Edward Hart
Location: Biotech-Chem Library
REM-1A55
Phone: 571-272-2512

edward.hart@uspto.gov

Search Notes

Examiner Fredman,

Here are the results of the search you requested.

Please feel free to contact me if you have any questions.

Edward Hart

STIC-Biotech/ChemLib

168245

mg

From: Fredman, Jeffrey
Sent: Tuesday, October 11, 2005 10:23 AM
To: STIC-Biotech/ChemLib
Subject: 09/744,097

RECEIVED
OCT 11 2005
STIC-BIOTECH/CHM LIB
(STIC)

Please search SEQ ID NO: 76 in nucleic acid databases.

Thanks,

Jeffrey Fredman
Art Unit 1637
Remsen Building 2C89
(571)272-0742

2C18

Searcher: _____
Searcher Phone: _____
Date Searcher Picked up: 10/12/05
Date completed: 10/14/05
Searcher Prep Time: _____
Online Time: _____

Type of Search
NA# 1 AA# _____
S/L: _____ Oligomer: _____
Encode/Transl: _____
Structure #: _____ Text: _____
Inventor: _____ Litigation: _____

Vendors and cost where applicable
STN: _____
DIALOG: _____
QUESTEL/ORBIT: _____
LEXIS/NEXIS: _____
SEQUENCE SYSTEM: _____
WWW/Internet: _____
Other (Specify): _____

GenCore version 5.1.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 13, 2005, 17:46:38 ; Search time 1479 Seconds
(without alignments)
884.578 Million cell updates/sec

Title: US-09-744-097A-76
Perfect score: 27
Sequence: 1 gtacctagctaccctaggtctaggc 27

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*

- 1: gb.ba.*
- 2: gb.htg.*
- 3: gb.in.*
- 4: gb.om.*
- 5: gb.ov.*
- 6: gb.pat.*
- 7: gb.ph.*
- 8: gb.pl.*
- 9: gb.pr.*
- 10: gb.ro.*
- 11: gb.sts.*
- 12: gb.sy.*
- 13: gb.un.*
- 14: gb.vl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20.6	76.3	135412	2	AC148279
2	20.6	76.3	156661	2	AC148358
3	20.6	76.3	278310	2	AC127955
C 4	20.6	76.3	285193	2	AC111242
C 5	20.6	76.3	293962	2	AC112303
6	20.4	75.6	189946	10	AC127236
C 7	20.4	75.6	194613	2	AC116089
C 8	20.2	74.8	124104	8	AC134931
9	19.8	73.3	119743	2	AC134515
10	19.8	73.3	211465	2	AC103070
11	19.8	73.3	212559	2	AC095896
12	19.6	72.6	857	4	SSAPOA11
13	19.6	72.6	5551	10	AK173314
C 14	19.6	72.6	207223	10	AL732521
15	19.6	72.6	212481	10	AC126959
16	19.6	72.6	252851	2	AC135094
C 17	19.2	71.1	276137	2	AC131170
18	19.2	71.1	91977	10	AL732588
19	19.2	71.1	116926	10	AL808012

20	19.2	71.1	147467	2	AC118573
C 21	19.2	71.1	186449	2	AC147134
22	19.2	71.1	188892	2	AC091327
C 23	19.2	71.1	237378	2	AC087038
24	19	70.4	110000	1	AE017180.27
25	19	70.4	150222	2	AC120016
26	19	70.4	164530	10	AL596104
27	19	70.4	187514	2	AC109240
C 28	19	70.4	203946	2	AC069465
C 29	19	70.4	233852	2	AC108549
C 30	19	70.4	235241	10	AL928893
C 31	19	70.4	240264	2	AC107434
32	19	70.4	245134	2	AC126639
33	19	70.4	248917	2	AC111455
C 34	19	70.4	250579	2	AC126843
35	19	70.4	323223	2	AC109504
C 36	18.8	69.6	81368	2	AC094244.4
C 37	18.8	69.6	103428	2	AC096436.6
C 38	18.8	69.6	104113	10	AL928640
C 39	18.8	69.6	110000	2	AC102028.3
40	18.8	69.6	142937	2	AC139916
C 41	18.8	69.6	148320	10	AL669849
C 42	18.8	69.6	154685	10	AL606511
C 43	18.8	69.6	157088	10	AC127292
C 44	18.8	69.6	171936	10	AL929035
C 45	18.8	69.6	173598	10	AL672194

ALIGNMENTS

AC148279 135412 bp DNA linear HTG 15-APR-2004
Sorex araneus clone SA_Ba-546N4, WORKING DRAFT SEQUENCE, 7 ordered pieces.

AC148279
AC148279.2 GI:46391177

HTG; HTGS PHASE2; HTGS_DRAFT.

Sorex araneus (European shrew)

Sorex araneus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Insectivora; Soricidae; Soricinae; Sorex.

1 (bases 1 to 135412)

Antonnellis, A., Ayele, K., Benjamin, B., Blakesley, R.W.

Bouffard, G.A., Brinkley, C., Brooks, S., Chu, G., Coleman, B.,

Coleman, H., Daki, N., Engle, J., Granite, S., Guan, X., Gupta, J.,

Haghighi, P., Han, J., Hansen, N., Ho, S.-L., Hu, P., Hurie, B.,

Idol, J.R., Jones, C., Karlins, E., Kim, H., Kwong, P., Laric, P.,

Larson, S., Lee-Lin, S.-O., Legaspi, R., Maduro, O.L., Maduro, V.B.,

Margulies, E.H., Masiello, C., Maskeri, B., McDowell, J.,

Mullikin, J.C., Paguirigan, C., Portnoy, M.E., Prased, A., Puri, O.,

Reddix-Dugue, N., Schandler, K., Schueler, M.G., Shah, K., Sison, C.,

Stanthrop, S., Thomas, J.W., Thomas, P.J., Tsipouri, V., Vogt, J.L.,

Wetherby, K.D., Young, A. and Green, E.D.

NISC Comparative Sequencing Initiative

Unpublished

2 (bases 1 to 135412)

Green, E.D.

Direct Submission

Submitted (19-FEB-2004) NIH Intramural Sequencing Center, 8717

Grovenmont Circle, Gaithersburg, MD 20877, USA

3 (bases 1 to 135412)

Green, E.D.

Direct Submission

Submitted (15-APR-2004) NIH Intramural Sequencing Center, 8717

Grovenmont Circle, Gaithersburg, MD 20877, USA

On Apr 15, 2004 this sequence version replaced gi:42627935.

----- Genome Center

Center: NIH Intramural Sequencing Center

Center code: NISC

Web site: <http://www.nisc.nih.gov>

Contact: nisc_zoo@nhgri.nih.gov

----- Project Information
 Center project name: gbi
 Center clone name: 546N04

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicated order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones), alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8X average coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

----- Summary Statistics

Sequencing vector: plasmid; n/a; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 134185 bases at least Q40
 Consensus quality: 134570 bases at least Q30
 Consensus quality: 134696 bases at least Q20
 Insert size: 118000; agarose-fp
 Insert size: 134812; sum-of-contigs
 Quality coverage: 11.34x in Q20 bases; agarose-fp
 Quality coverage: 9.93x in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently consists of 7 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.

* This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.

1 28824: contig of 28824 bp in length
 28825 28924: gap of unknown length
 28925 35906: contig of 6982 bp in length
 35907 36006: gap of unknown length
 36007 61654: contig of 25648 bp in length
 61655 61754: gap of unknown length
 61755 68663: contig of 6909 bp in length
 68664 68763: gap of unknown length
 68764 126764: contig of 58001 bp in length
 126765 126865: gap of unknown length
 133235: contig of 6371 bp in length
 133236 133335: gap of unknown length
 133336 133412: contig of 2077 bp in length.

FEATURES

source

1. 135412
 /organism="Sorex araneus"
 /mol_type="genomic DNA"
 /db_xref="taxon:42254"
 /clone="SA_Ba-546N4"
 /clone_lib="SA_Ba"

misc_feature

1. 60485
 /note="clone overlaps with GenBank Accession Number AC148351 clone SA_Ba-157G15 (center project name gbff)"

misc_feature

1. 28824
 /note="assembly_fragment
 clone_end:T7
 vector_side:left"

misc_feature

28925. 35906

misc_feature

36007. 61654

misc_feature

61755. 68663

misc_feature

68764. 126764

misc_feature

69707. 135412

/note="clone overlaps with GenBank Accession Number AC148358 clone SA_Ba-621C10 (center project name gbh)"
 126865. 133235
 /note="assembly_fragment"
 13336. 135412
 /note="assembly_fragment
 clone_end:SP6
 vector_side:right"

ORIGIN

Query Match 76.3%; Score 20.6; DB 2; Length 135412;

Best Local Similarity 85.2%; Pred. No. 20;

Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 GTACCTAGTACCCCTAGCTAGGC 27

Db 109081 GTACCTAAGTACCCCTAGATTAGGC 109107

RESULT 2

AC148358

LOCUS

DEFINITION Sorex araneus clone SA_Ba-621C10, WORKING DRAFT SEQUENCE, 11 ordered pieces.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

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coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

----- Summary Statistics

Sequencing vector: plasmid; n/a; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 154585 bases at least Q40
 Consensus quality: 155145 bases at least Q30
 Consensus quality: 155506 bases at least Q20
 Insert size: 107000; agarose-fp
 Insert size: 155661; sum-of-contigs
 Quality coverage: 12.02x in Q20 bases; agarose-fp
 Quality coverage: 8.26x in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.

* This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.

* 1 41408: contig of 41408 bp in length
 * 41409: gap of unknown length
 * 41509: contig of 2126 bp in length
 * 43635: gap of unknown length
 * 43735: contig of 13527 bp in length
 * 57262: gap of unknown length
 * 57361: contig of 6500 bp in length
 * 63861: gap of unknown length
 * 63862: contig of 33808 bp in length
 * 63962: gap of unknown length
 * 97770: contig of 10730 bp in length
 * 97870: contig of 10730 bp in length
 * 108600: gap of unknown length
 * 108700: contig of 20478 bp in length
 * 129178: gap of unknown length
 * 129278: contig of 5413 bp in length
 * 134691: gap of unknown length
 * 134791: contig of 7141 bp in length
 * 141931: gap of unknown length
 * 142032: contig of 9880 bp in length
 * 151912: gap of unknown length
 * 152012: contig of 4650 bp in length.

FEATURES

source

1. 155661
 /organism="Sorex araneus"
 /mol_type="genomic DNA"
 /db_xref="taxon:42254"
 /clone="SA_Ba-621C10"
 /clone_lib="SA_Ba"

1. 67579

/note="clone overlaps with GenBank Accession Number AC148279, clone SA_Ba-546N4 (center project name gbi)"

1. 41408
 /note="assembly_fragment
 clone_end:SP6
 vector_side:left"

41509. 43634

/note="assembly_fragment"
 43735. 57261
 /note="assembly_fragment"

57362. 63861

/note="assembly_fragment"
 63962. 97769
 /note="assembly_fragment"

97870. 108599

/note="assembly_fragment"
 108700. 129177
 /note="assembly_fragment"

129278. 134690

misc_feature
 /note="assembly_fragment"
 129878. 155661
 /note="clone overlaps with GenBank Accession Number AC148355, clone SA_Ba-408O18 (center project name gga)"
 misc_feature
 134791. 141931
 /note="assembly_fragment"
 misc_feature
 142032. 151911
 /note="assembly_fragment"
 misc_feature
 152012. 155661
 /note="assembly_fragment
 clone_end:T7
 vector_side:right"

ORIGIN

Query Match 76.3%; Score 20.6; DB 2; Length 155661;
 Best Local Similarity 85.2%; Fred. No. 20;
 Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
 Oy 1 GTAGCTAGCTACCCCTAGGTCTAGGC 27
 Db 39469 GTACCTACTACCCCTAGATTAGGC 39495

RESULT 3

AC127955

LOCUS

DEFINITION AC127955 278310 bp DNA linear HTG 19-SEP-2002
 Rattus norvegicus clone CH230-270L18, *** SEQUENCING IN PROGRESS
 ***, 6 unordered pieces.

AC127955

ACCESSION

AC127955.2 GI:23195969

VERSION

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.

KEYWORDS

Rattus norvegicus (Norway rat)

SOURCE

ORGANISM

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus.

1 (bases 1 to 278310)

Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,

Allen, C., Allen, H., Alebrooks, S., Amin, A., Anguiano, D.,

Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,

Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,

Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,

Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,

Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,

Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,

Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,

Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,

Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,

Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,

Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G.,

Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,

Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, I., Garza, M.,

Gebrgeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, I., Guerra, W.,

Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K.,

Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J.,

Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hogues, M.,

Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A.,

Jackson, S., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jollivet, A.,

Karpathy, S., Kelly, S., Khan, Z., King, L., Kovar, C.,

Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,

Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,

Lorensuhewa, L., Loulseghe, H., Lozano, R. J., Lu, X., Ma, J.,

Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,

Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,

Mawhney, S., McLeod, M. P., McNeill, T. Z., Meenen, E.,

Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,

Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,

Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,

Nwaokemele, O., Okwuon, G., Olarnpunsagoon, A., Pal, S., Parks, K.,

Pasternak, S., Paul, H., Perez, A., Perez, B., Pfankuch, C.,

Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L. L.,

Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R.,

Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,

Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sison, I., Sitter, C.D., Smajls, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorrelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmami, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willison, R., Wleciyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, V., Yu, F., Zhang, J., Zhou, X., Zhou, X., Zhao, S., Dunn, D., von Niederhausen, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

TITLE
JOURNAL
REFERENCE
AUTHORS
JOURNAL

2 (bases 1 to 278310)
Worley K.C.
Direct Submission
Submitted (19-JUL-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 278310)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (19-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

REFERENCE
AUTHORS
JOURNAL

COMMENT

The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). As a result, the sequence may extend beyond the ends of the clone and there may be contigs that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: KBR
Center clone name: CH230-270L18
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 159405 bases at least Q40
Consensus quality: 163261 bases at least Q30
Consensus quality: 165346 bases at least Q20
Estimated insert size: 206482; sum-of-contigs estimation
Quality coverage: 3x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently consists of 6 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

* 1 197344: contig of 197344 bp in length
* 197445: gap of unknown length
* 197445 266948: contig of 69504 bp in length
* 266949 267048: gap of unknown length
* 267049 269714: contig of 2666 bp in length
* 269715 269814: gap of unknown length
* 269815 271799: contig of 1985 bp in length
* 271800 271899: gap of unknown length
* 271900 273966: contig of 2067 bp in length
* 273967 274066: gap of unknown length
* 274067 278310: contig of 4244 bp in length.

Location/Qualifiers

source
1. 278310
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/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-270L18"
misc_feature
1. 2376
/note="wgs_contig"
153418..156139
misc_feature
/note="wgs_contig"
211774..212782
misc_feature
/note="wgs_contig"
ORIGIN

Query Match 76.3%; Score 20.6; DB 2: Length 278310;
Best Local Similarity 85.2%; Pred. No. 19;
Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 1 GTACCTAGCTACCCCTAGGCTAGGC 27
Db 248352 GTAGCCAGGTACCTAGGCTAGC 248378

RESULT 4
AC111242/c
LOCUS
AC111242 Rattus norvegicus clone CH230-232H4, linear HTG 13-MAY-2003
DEFINITION
***, 11 unordered pieces.
AC111242
AC111242.4 GI:30578456
VERSION
HTG: HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
KEYWORDS
Rattus norvegicus (Norway rat)
SOURCE
Rattus norvegicus
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

REFERENCE
1 (bases 1 to 285193)
Muzny, D., Maric, E., Metzker, M., Lee, A., Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Ayagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denison, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, J., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorenschew, L., Loulseghe, H., Lozada, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhinney, S., Mcleod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokeme, O., Okwono, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, I., Pfannkuch, C., Plopper, F., Polindexter, A., Popovic, D., Primus, E., Pu, L., Reigh, R., Puzo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J.,

Sanders, W., Saverly, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Taber, P., Taylor, C., Taylor, T., Thera, N., Thomas, S., Tingey, A., Trejos, Z., Umani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

TITLE JOURNAL

REFERENCE AUTHORS

TITLE JOURNAL

REFERENCE AUTHORS

TITLE JOURNAL

REFERENCE AUTHORS

TITLE JOURNAL

COMMENT

2 (bases 1 to 285193)
Worley, K.C.

Submitted (19-FEB-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 285193)
Rat Genome Sequencing Consortium.

Submitted (13-MAY-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On May 13, 2003 this sequence version replaced gi:23604121.

The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu

Project Information
Center project name: GWE
Center clone name: CH230-232H4

Assembly program: Atlas 3.0;
Consensus quality: 242831 bases at least Q40
Consensus quality: 249455 bases at least Q30
Consensus quality: 253771 bases at least Q20

Estimated insert size: 272024; sum-of-contigs estimation
Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

NOTE: Estimated insert size may differ from sequence length (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)

NOTE: This sequence may represent more than one clone.

NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.

This record will be updated with the finished sequence, as soon as it is available and the accession number will be preserved.

1 5183: contig of 5183 bp in length
5184 5283: gap of unknown length
5284 17822: contig of 12539 bp in length
17823 116644: contig of 98722 bp in length
116645 157368: contig of 40624 bp in length

* 157369 157468: gap of unknown length
157469 247903: contig of 90435 bp in length
247904 248003: gap of unknown length
248004 261891: contig of 13888 bp in length
261892 261991: gap of unknown length
261992 276510: contig of 14519 bp in length
276511 276610: gap of unknown length
276611 277878: contig of 1268 bp in length
277879 277978: gap of unknown length
277979 279325: contig of 1347 bp in length
279326 279425: gap of unknown length
279426 282175: contig of 2750 bp in length
282176 282275: gap of unknown length
282276 285193: contig of 2918 bp in length.

FEATURES source

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

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misc_feature

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/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-232H4"

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/note="wgs_end_extension"
clone_end:T7

complement(7161..8031)
/note="clone_boundary"
clone_end:T7

end_sequence:BZ092926"
16581..17822
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30287..34910
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34961..39614
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40916..42387
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58872..59764
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113101..116644
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136932..138724
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155552..157368
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clone_end:Sp6"
257732..259507
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268633..274031
/note="wgs_end_extension"

clone_end:Sp6"

Query Match 76.3%; Score 20.6; DB 2; Length 285193;
Best Local Similarity 85.2%; Pred. No. 19;
Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 GTAGCTAGCTACCCCTAGGCTTAGGC 27
|||||

Db 114184 GTAGCAAGCTACCTCTAGGTGTAGAC 114158
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RESULT 5
AC112303/c AC112303 293962 bp DNA linear HTG 15-NOV-2002

LOCUS Rattus norvegicus clone CH230-208p20, *** SEQUENCING IN PROGRESS

DEFINITION *** 8 unordered pieces.

ACCESSION AC112303

```

VERSION
KEYWORDS
SOURCE
ORGANISM

AC12303.4 GI:24635626
HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE
AUTHORS
Muzny, D., Marie, Metzger, M., Lee, Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, P.,
Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, J., Burch, P., Burrell, K., Calderon, E.,
Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, J., Chu, J.,
Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,
Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G.,
Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,
Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,
Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,
Gunnarane, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K.,
Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J.,
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Hollins, B., Howells, S., Huiyk, S., Hume, J., Idlebird, D., Jackson, A.,
Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,
Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,
Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,
Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
Lorensheva, L., Louisedge, H., Lozada, R.J., Lu, X., Ma, J.,
Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,
Mangum, B., Mapa, P., Martin, K., Martin, R., Martinez, E.,
Mahoney, S., McLeod, M.P., McNeill, T.Z., Meenen, E.,
Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,
Morgan, M., Morris, K., Morris, S., Munitasa, M., Murphy, M., Nair, L.,
Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,
Nwakoleme, O., Okwuonu, G., Olarunsaogoon, A., Pal, S., Parks, K.,
Pasternak, S., Paul, H., Perez, A., Perez, L., Pfankuch, L.,
Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.,
Puafo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,
Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,
Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.O.,
Sanders, W., Savery, G., Scherer, S., Scott, G., Shatman, S., Shen, H.,
Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D.,
Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J.,
Steinle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,
Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Umani, K.,
Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,
Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,
Williams, G., Willson, R., Wleczky, R., Wooden, H., Worley, K.,
Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,
Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,
Weinstock, G. and Gibbs, R.A.
Direct Submission
Unpublished
2 (bases 1 to 293962)
Worley, K.C.

TITLE
JOURNAL
REFERENCE
AUTHORS

Direct Submission
Submitted (21-FEB-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 293962)

REFERENCE
AUTHORS
JOURNAL

Rat Genome Sequencing Consortium.
Submitted (15-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Nov 6, 2002 this sequence version replaced gi:23603916.
The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GMYH
Center clone name: CH230-208P20
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 206616 bases at least Q40
Consensus quality: 209134 bases at least Q30
Consensus quality: 211305 bases at least Q20
Estimated insert size: 210335; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 27340: contig of 27340 bp in length
* 27341 27440: gap of unknown length
* 27441 51601: contig of 24161 bp in length
* 51602 51701: gap of unknown length
* 51702 268441: contig of 216740 bp in length
* 268442 270645: gap of unknown length
* 270646 270745: gap of unknown length
* 270746 272239: contig of 1493 bp in length
* 272239 27338: gap of unknown length
* 27338 275355: contig of 3016 bp in length
* 275355 275454: gap of unknown length
* 275455 281008: contig of 5554 bp in length
* 281009 293962: contig of 12854 bp in length.
* 293962
Location/Qualifiers
1. 293962
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-208P20"
6200..7062
/clone="clone_boundary"
/clone_end:Sp6
misc_feature
end sequence:RWBKP94TVB"
51702..53500
/clone="wgs_contig"
53551..54607
/clone="wgs_contig"
56203..57503
/clone="wgs_contig"
174983..176056
/clone="wgs_contig"
208600..209401
/clone="clone_boundary"

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clone_end:T7
site:
end sequence:RWBKP94TJB"
misc_feature      264125..265539
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clone_end:T7"
misc_feature      267070..268441
                  /note="wgs_end_extension
clone_end:T7"

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ORIGIN

Query Match 76.3%; Score 20.6; DB 2; Length 293962;

Best Local Similarity 85.2%; Pred. No. 19;

Matches 23; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 1 GTAGCTAGTACCCCTAGTCTAGGC 27

Db 256250 GTAGCCAGCTACCTTAGTGTAGAC 256224

RESULT 6

AC127236

LOCUS AC127236 188946 bp DNA linear ROD 27-NOV-2003
DEFINITION Mus musculus BAC clone RP24-351L1 from chromosome 18, complete
sequence.

ACCESSION

VERSION AC127236.3 GI:37361086

KEYWORDS

HTG.

SOURCE

ORGANISM Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 188946)

Swearngen-Shahid, S.

The sequence of Mus musculus BAC clone RP24-351L1

Unpublished (2001)

2 (bases 1 to 188946)

Wilson, R.

Sequencing of Mus musculus

Unpublished (2001)

3 (bases 1 to 188946)

McPherson, J.D. and Waterston, R.H.

Direct Submission

Submitted (14-JUL-2002) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

4 (bases 1 to 188946)

McPherson, J.D. and Waterston, R.H.

Direct Submission

Submitted (25-SEP-2002) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

5 (bases 1 to 188946)

Wilson, R.K.

Direct Submission

Submitted (02-OCT-2003) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

6 (bases 1 to 188946)

Wilson, R.

Direct Submission

Submitted (27-NOV-2003) Department of Genetics, Washington

University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

On Oct 2, 2003 this sequence version replaced gi:23308121.

----- Genome Center

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: <http://genome.wustl.edu>

Contact: submissions@watson.wustl.edu

----- Summary Statistics

Center project name: M_BB0351L01

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap

between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. Wes Warren,
Department of Genetics, Washington University, St. Louis MO. For
additional information about the map position of this sequence, see
<http://genome.wustl.edu>

SOURCE INFORMATION:

The RPCI-24 BAC library has been constructed by Pieter de Jong and
coworkers (<http://www.chori.org>) from male C57BL/6J mouse spleen
and/or brain genomic DNA. The clone and detailed information can be
obtained from Pieter de Jong and coworkers at <http://www.chori.org>

NEIGHBORING SEQUENCE INFORMATION:

This sequence is the entire insert of the clone. This clone is
overlapped by AC108434.

FEATURES

source

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Location/Qualifiers

/organism="Mus musculus"

/mol_type="genomic DNA"

/db_xref="taxon:10090"

/chromosome="18"

/map="18"

/clone="RP24-351L1"

/clone_lib="RPCI-24"

repeat_region 773..874

/rpt_family="Alu"

repeat_region 920..1078

/rpt_family="Alu"

repeat_region 1428..1643

/rpt_family="B2"

repeat_region 1675..1845

/rpt_family="Alu"

repeat_region 2684..2873

/rpt_family="B2"

repeat_region 3164..3303

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repeat_region 3577..3670

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repeat_region 3688..3892

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repeat_region 5876..6150

/rpt_family="B4"

repeat_region 6252..6351

/rpt_family="Alu"

repeat_region 6452..6680

/rpt_family="B4"

repeat_region 6545..6681

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repeat_region 6698..6951

/rpt_family="B4"

repeat_region 7324..7516

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repeat_region 10777..10894

/rpt_family="B4"

repeat_region 11023..11190

/rpt_family="B4"

repeat_region 11031..11227

/rpt_family="B2"

repeat_region 11303..11433

/rpt_family="Alu"

repeat_region 11547..11724

/rpt_family="B2"

repeat_region 11736..12221

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Best Local Similarity 95.5%; Pred. No. 24;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 CTAGCTACCCCTAGGCTCTAGGC 27
||| ||||| ||||| ||||| |||||
Db 132397 CTAACCTACCCCTAGGCTCTAGGC 132418

RESULT 7
AC116089/c
LOCUS AC116089 194613 bp DNA linear HTG 20-NOV-2002
DEFINITION Rattus norvegicus clone CH210-344011, WORKING DRAFT SEQUENCE, 2
unordered pieces.
AC116089
ACCESSION AC116089, 6 GI:25138296
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE 1 (bases 1 to 194613)
AUTHORS Kuzny, D., Marie, Metzker, M., Lee, Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Aisbrooks, S., Amin, A., Angiano, D.,
Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, B., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
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Georgiev, E., Geer, K., Gill, R., Grady, M., Guarra, W., Guevara, W.,
Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K.,
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Karpach, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,
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Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
Lorensu, L., Lorensu, L., Lozada, R., Lu, X., Lu, X., Ma, J.,
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Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,
Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E.,
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Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C.,
Plopper, F., Polindexter, A., Popovic, D., Primus, E., Pu, L., L.,
Puato, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,
Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,
Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S., J.,
Sanders, W., Savary, G., Scheer, S., Scott, G., Scott, S., Shen, H.,
Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D.,
Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J.,
Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,
Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K.,
Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,
Wang, O., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,
Williams, G., Willson, R., Wleczky, R., Woodson, H., Worley, K.,
Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, V., Yoon, V.,
Yoon, V.,

Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.
 Direct Submission
 Unpublished
 2 (bases 1 to 194613)
 Worley, K.C.
 Direct Submission
 Submitted (24-MAR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 194613)
 Rat Genome Sequencing Consortium.
 Direct Submission
 Submitted (20-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 On Nov 20, 2002 this sequence version replaced gi:23194943.
 The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
 Center: Baylor College of Medicine
 Center code: BCM
 Web site: <http://www.hgsc.bcm.tmc.edu/>
 Contact: hgsc-help@bcm.tmc.edu
 ----- Project Information
 Center project name: GTCO
 Center clone name: CH230-344011
 ----- Summary Statistics
 Assembly program: Phrap; version 0.990329
 Consensus quality: 178038 bases at least Q40
 Consensus quality: 179671 bases at least Q30
 Consensus quality: 180674 bases at least Q20
 Estimated insert size: 180448; sum-of-contigs estimation
 Quality coverage: 8x in Q20 bases; sum-of-contigs estimation

 * NOTE: Estimated insert size may differ from sequence length
 * (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
 * NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

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 DB 80840 CTAGCTACCCCTAGGCTTAGGC 80819
 RESULT 8
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 LOCUS
 DEFINITION Oryza sativa (japonica cultivar-group) chromosome 5 clone
 OSJNB007911, complete sequence.
 ACCESSION AC134931
 VERSION AC134931.2 GI:37360982
 KEYWORDS HTG.
 SOURCE Oryza sativa (japonica cultivar-group)
 ORGANISM Oryza sativa (japonica cultivar-group)
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzaceae; Oryza.
 1 (bases 1 to 124104)
 Chow, T.-Y., Hsing, Y.-I.C., Chen, C.-S., Chen, H.-H., Liu, S.-M.,
 Chao, Y.-T., Chang, S.-J., Chen, H.-C., Chen, S.-K., Chen, T.-R.,
 Chen, Y.-L., Cheng, C.-H., Chung, C.-I., Han, S.-Y., Hsiao, S.-H.,
 Hsiung, J.-N., Hsu, C.-H., Huang, J.-J., Kau, P.-I., Lee, M.-C.,
 Leu, H.-L., Li, Y.-F., Lin, S.-J., Lin, Y.-C., Wu, S.-W., Yu, C.-Y.,
 Yu, S.-W., Wu, H.-P. and Shaw, J.-F.
 Oryza sativa BAC OSJNB007911 genomic sequence
 Unpublished
 2 (bases 1 to 124104)
 Chow, T.-Y. and Hsing, Y.-I.C.
 Direct Submission
 Submitted (03-OCT-2002) Institute of Botany, Academia Sinica, 128,
 Section 2, Academia Road, Nankang, Taipei 11529, Taiwan
 3 (bases 1 to 124104)
 Chow, T.-Y.
 Direct Submission
 Submitted (02-OCT-2003) Institute of Botany, Academia Sinica, 128,
 Section 2, Academia Road, Nankang, Taipei 11529, Taiwan
 4 (bases 1 to 124104)
 Chow, T.-Y. and Hsing, Y.-I.C.
 Direct Submission
 Submitted (02-JUN-2004) Institute of Botany, Academia Sinica, 128,
 Section 2, Academia Road, Nankang, Taipei 11529, Taiwan
 On Oct 2, 2003 this sequence version replaced gi:23477780.
 The orientation of the sequence is from Sp6 to T7 of the BAC clone.
 Genes were predicated from the integrated results of the following:
 BLASTN2.0, BLASTX2.0, GENSCAN (Chris Burge,
<http://genes.mit.edu/GENSCAN.html>), Egenes
<http://www.softberry.com/>), GlimmerR
<http://www.tigr.org/softlab/glimmer/glimmer.html>), TWINSKAN
<http://genes.cs.wustl.edu/>) and Geneslicer
<http://www.tigr.org/tdb/Geneslicer/index.shtml>). The sequence was
 searched against the Swiss-Prot+TrEMBL protein database, the NCBI
 Plant EST database, the TIGR Rice Gene Index and the rice
 full-length cDNA database (KOME,
<http://cdna01.dna.affrc.go.jp/cDNA/>). Annotated genes are named to
 indicate the level of evidence for their annotation. Genes with

similarity to other proteins are named after the database hits. Genes without significant peptide similarity but with EST similarity are named as unknown proteins. Genes without protein or EST similarity, that are predicted by more than two gene prediction programs over most of their length are annotated as hypothetical proteins. This clone overlaps with P0008A07 (accession # AC079021) and P0583F12 (accession # AC129720).

FEATURES

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RESULT 9
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LOCUS      119743 bp      DNA      linear      HTG 27-SEP-2002
DEFINITION Rattus norvegicus clone CH230-1016, *** SEQUENCING IN PROGRESS ***,
63 unordered pieces.
AC134515
AC134515.1 GI:23334689
VERSION    HTG; HTGS_PHASE1
KEYWORDS   Rattus norvegicus
SOURCE     Rattus norvegicus
ORGANISM   Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 119743)
Muzny, D., Marle, Metzker, M., Lee, S., Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Ayoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
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Song, X., Z., Sorrell, R., Sosa, J., Steimle, M., Strong, R., Sutton, A.,
Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S.,
Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D.,
Walldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J.,
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TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S.,
Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X.,
Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D. R.,
Holt, R. A., Smith, H. O., Weinstock, G. and Gibbs, R. A.
Direct Submission
Unpublished
2 (bases 1 to 119743)
Rat Genome Sequencing Consortium
Direct Submission
Submitted (27-SEP-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GPSS
Center clone name: CH230-1016
----- Summary Statistics
Sequencing vector: Plasmid;
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 65609 bases at least Q40
Consensus quality: 70531 bases at least Q30
Consensus quality: 73895 bases at least Q20

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 63 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence.
* As soon as it is available and the accession number will
* be preserved.

* 1 1021: contig of 1021 bp in length
* 1022 1121: gap of unknown length
* 1122 2410: contig of 1289 bp in length
* 2411 2510: gap of unknown length
* 2511 3615: contig of 1105 bp in length
* 3616 3715: gap of unknown length
* 3716 5208: contig of 1493 bp in length
* 5209 5308: gap of unknown length
* 5309 6348: contig of 1040 bp in length
* 6349 6449: gap of unknown length
* 6449 7557: contig of 1108 bp in length
* 7557 7656: gap of unknown length
* 7657 8690: contig of 1034 bp in length
* 8691 8790: gap of unknown length
* 8791 9582: contig of 1192 bp in length
* 9583 10082: gap of unknown length
* 10083 11108: contig of 1026 bp in length
* 11089 11209: gap of unknown length
* 11209 12381: contig of 1173 bp in length
* 12382 12482: gap of unknown length
* 12482 13945: contig of 1463 bp in length
* 13945 14044: gap of unknown length
* 14045 15293: contig of 1249 bp in length
* 15294 15393: gap of unknown length
* 15394 16836: contig of 1443 bp in length
* 16837 16936: gap of unknown length
* 16937 18367: contig of 1331 bp in length
* 18368 18367: gap of unknown length
* 18368 19813: contig of 1446 bp in length
* 19814 19913: gap of unknown length
* 19914 21224: contig of 1311 bp in length
* 21225 21324: gap of unknown length
* 21325 22712: contig of 1388 bp in length
* 22713 22812: gap of unknown length
* 22813 23831: contig of 1019 bp in length

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 23932 25886: contig of 1955 bp in length
 25887 25986: gap of unknown length
 27306: contig of 1320 bp in length
 27406: gap of unknown length
 28973: contig of 1567 bp in length
 29073: gap of unknown length
 30417: contig of 1344 bp in length
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 31979: contig of 1462 bp in length
 32079: gap of unknown length
 33446: contig of 1367 bp in length
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 35631: contig of 2085 bp in length
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 37564: contig of 1833 bp in length
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 38768: contig of 1104 bp in length
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 39951: contig of 1083 bp in length
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 41348: contig of 1297 bp in length
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 42876: contig of 1428 bp in length
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 44226: contig of 1250 bp in length
 44326: gap of unknown length
 45971: contig of 1645 bp in length
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 48011: contig of 1940 bp in length
 48111: gap of unknown length
 49629: contig of 1518 bp in length
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 51576: contig of 1847 bp in length
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 53467: contig of 1791 bp in length
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 54758: contig of 1191 bp in length
 54858: gap of unknown length
 56642: contig of 1784 bp in length
 56742: gap of unknown length
 58752: contig of 2010 bp in length
 58852: gap of unknown length
 60328: contig of 1476 bp in length
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 62184: contig of 1756 bp in length
 62284: gap of unknown length
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 66273: gap of unknown length
 67696: contig of 1423 bp in length
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 71636: gap of unknown length
 73791: contig of 2155 bp in length
 73891: gap of unknown length
 76129: contig of 2238 bp in length
 76229: gap of unknown length
 78594: contig of 2365 bp in length
 78694: gap of unknown length
 80868: contig of 2174 bp in length
 80968: gap of unknown length
 82506: contig of 1538 bp in length
 82606: gap of unknown length
 85214: contig of 2608 bp in length
 85314: gap of unknown length
 87565: contig of 2251 bp in length
 87655: gap of unknown length

Query Match

73.3%; Score 19.8; DB 2; Length 119743;

Best Local Similarity 91.3%; Pred. No. 50;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 5 CCTAGTACCCCTAGGTCTAGGC 27
 DB 93195 CCTACTACCCCTAGGACTAGGC 93217
 RESULT 10
 AC103070
 LOCUS
 DEFINITION
 Rattus norvegicus clone CH230-17IN16, *** SEQUENCING IN PROGRESS
 AC103070 211465 bp DNA linear HTG 13-MAY-2003
 Rattus norvegicus clone CH230-17IN16, *** SEQUENCING IN PROGRESS
 AC103070 211465 bp DNA linear HTG 13-MAY-2003
 AC103070 6 GI:30579789
 HTG: HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
 Rattus norvegicus (Norway rat)
 SOURCE
 ORGANISM
 Rattus norvegicus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.
 1 (bases 1 to 211465)
 Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J.,
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
 Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
 Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
 Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
 Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
 Chacko, J., Chavez, D., Chen, R., Chen, Y., Chen, Y., Chu, J.,
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
 Davila, M. L., Davis, C., Davy-Carroil, L., De Andra, C., Dederich, D.,
 Delgado, O., Denson, S., Dey, C., Ding, Y., Dinh, H., Divya, K.,
 Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,
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 Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K.,
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 Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jollivet, A.,
 Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Liu, J.,
 Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,
 Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
 Lorensu, H., Loulseghe, H., Lozano, R. J., Lu, X., Ma, J.,
 Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,
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 Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J.,
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 Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Umani, K.,
 Valas, R., Vera, V., Villanasa, D., Waldron, L., Walker, B., Wang, J.,
 Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,
 Williams, G., Willson, R., Wlecyk, R., Woodson, H., Worley, K.,
 Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,
 Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
 Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O.,
 Weinstock, G. and Gibbs, R. A.
 Direct Submission
 TITLE
 JOURNAL
 Unpublished

REFERENCE

2 (bases 1 to 211465)

Worley, K.C.
Direct Submission
Submitted (24-NOV-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

REFERENCE

3 (bases 1 to 211465)

Submitted (13-MAY-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

COMMENT

On May 13, 2003 this sequence version replaced gi:23321751.
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GJDL

Center clone name: CH230-171N16

----- Summary Statistics

Assembly program: Atlas 3.0;

Consensus quality: 165184 bases at least Q40

Consensus quality: 169724 bases at least Q30

Consensus quality: 173497 bases at least Q20

Estimated insert size: 175743; sum-of-contigs estimation

Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

* 1 15062: contig of 15062 bp in length

* 15063 15162: gap of unknown length

* 15163 211465: contig of 196303 bp in length.

Location/Qualifiers

1. 211465

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-171N16"

9750. 11143

/notes="wgs contig"

31208. 33466

/note="wgs contig"

136133. 137193

/notes="wgs contig"

143435. 145324

/notes="wgs contig"

184822. 187065

/note="wgs contig"

189375. 189980

/notes="clone boundary

clone_end:Sp6

FEATURES

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misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

misc_feature

site:BCORI

end sequence: BH289014"

209353. 211465

/note="wgs_end_extension

clone_end:Sp6"

ORIGIN

Query Match 73.3%; Score 19.8; DB 2; Length 211465;

Best Local Similarity 91.3%; Pred. No. 48;

Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 CCTAGTACCCCTAGTCTAGGC 27

Db 22022 CCTAAGTACCCCTAGGACTAGGC 22044

RESULT 11

AC095896

LOCUS

DEFINITION

AC095896

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.

KEYWORDS

SOURCE

ORGANISM

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus.

1 (bases 1 to 212559)

Muzny, D. Marie, Metzker, M. Lee, Abramson, S., Adams, C., Alder, J.,

Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,

Anylebechi, V., Ayodeji, A., Ayodeji, M., Baca, E., Baden, H.,

Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,

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Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,

Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,

Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,

Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,

Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,

Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,

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Gunaratne, P., Haaland, N., Hamil, C., Hamilton, C., Hamilton, K.,

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Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,

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Lorenshewari, L., Louisedge, H., Lozada, R. J., Lu, X., Ma, J.,

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Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,

Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,

Nwackeleme, O., Okwuon, G., Olarnpunsagoon, A., Pal, S., Parks, K.,

Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C.,

Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L. L.,

Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R.,

Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,

Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J.,

Sanders, W., Savary, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,

Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C. D., Smays, D.,

Sneed, A., Sodergren, E., Song, X. Z., Sorelle, R., Sosa, J.,

Steinle, M., Strong, R., Sutton, A., Svatek, A., Taber, P., Taylor, C.,

Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K.,

Valas, R., Vera, V., Villalana, D., Waldron, L., Walker, B., Wang, J.,

Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F., Williams,G., Willson,R., Wlezyk,R., Wooden,H., Worley,K., Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V., Yu,F., Zhang,J., Zhou,X., Zhou,X., Zhao,S., Dunn,D., von Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O., Weinstock,G. and Gibbs,R.A.

Direct Submission
Unpublished
2 (bases 1 to 212559)
Worley,K.C.

Direct Submission
Submitted (17-SEP-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 212559)
Rat Genome Sequencing Consortium.

Direct Submission
Submitted (13-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On Nov 13, 2002 this sequence version replaced gi:23269609. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu

----- Project Information
Center project name: GDUL
Center clone name: CH230-10N20
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 172337 bases at least Q40
Consensus quality: 177428 bases at least Q30
Consensus quality: 180480 bases at least Q20
Estimated insert size: 178337; sum-of-contigs estimation
Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently consists of 10 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 45798: contig of 45798 bp in length
* 45799 45898: gap of unknown length
* 45899 146288: contig of 100370 bp in length
* 146289 146368: gap of unknown length
* 146369 150042: contig of 3674 bp in length
* 150043 150142: gap of unknown length
* 150143 150143: contig of 12189 bp in length
* 162332 162431: gap of unknown length
* 162432 199327: contig of 36896 bp in length
* 199328 199427: gap of unknown length
* 199428 200818: contig of 1391 bp in length
* 200819 200918: gap of unknown length
* 200919 202031: contig of 1113 bp in length
* 202032 202131: gap of unknown length

* 202132 203608: contig of 1477 bp in length
* 203609 203708: gap of unknown length
* 203709 206315: contig of 2607 bp in length
* 206316 206415: gap of unknown length
* 206416 212559: contig of 6144 bp in length.

FEATURES
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/db_xref="taxon:10116"
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misc_feature
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ORIGIN
Query Match 73.3%; Score 19.8; DB 2; Length 212559;
Best Local Similarity 91.3%; Pred. No. 48;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 CCTAGCTACCCCTAGGTCTAGGC 27
Db 90486 CCTACTATCCCTAGGTCTAGGC 90508

RESULT 12
SSAPOA11
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
source
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/organism="Sus scrofa"
/mol_type="genomic DNA"
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70..75
CAAT_signal
TATA_signal
Gene
mRNA
exon
intron

SSAPOA11
S.scrofa gene for apolipoprotein A1, exons 1-3.
X69478.1 GI:1887
apolipoprotein A-I; lecithin cholesterol acyltransferase cofactor;
lipid binding.
Sus scrofa (pig)
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
Mockel,B., Zinke,H., Flach,R., Weiss,B., Weiler-Guttler,H. and Gassen,H.G.
Expression of apolipoprotein A-I in porcine brain endothelium in vitro
J. Neurochem. 62 (2), 788-798 (1994)
94125128
8294940
2 (bases 1 to 857)
Moeckel,B.
Direct Submission
Submitted (27-NOV-1992) B. Moeckel, Inst. fuer Biochemie, Prof. Gassen, Technische Hochschule Darmstadt, Petersenstr. 22, 6100 Darmstadt, FRG
Location/Qualifiers
1..857
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/mol_type="genomic DNA"
/db_xref="taxon:9823"
/clone="PCR products"
/tissue type="liver"
/dev stage="adult"
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179..199
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200..368


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369..431
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/ number=3
766..857
/ gene="APOA1"
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ORIGIN

Query Match 72.6%; Score 19.6; DB 4; Length 857;
 Best Local Similarity 84.6%; Pred. No. 90;
 Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGTACCCCTAGGCTTAGGC 27
 Db 224 TACCCTAGTTCCTCCCGAGCTTAGGC 249

RESULT 13

AK173314 5551 bp mRNA linear ROD 28-JUL-2004
 LOCUS Mus musculus mRNA for mKIAA1973 protein.
 DEFINITION

ACCESSION AK173314

VERSION AK173314.1 GI:50511212

KEYWORDS FLI CDNA.

SOURCE Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

REFERENCE

AUTHORS

Okazaki,N., Kikuno,R.F., Ohara,T., Inamoto,S., Koseki,H.,
 Hiraoka,S., Saga,Y., Seino,S., Nishimura,M., Kaisho,T., Hoshino,K.,
 Kitamura,H., Nagase,T., Ohara,O. and Koga,H.
 Prediction of the Coding Nucleotide Sequences of KIAA
 Gene: IV. The Complete Nucleotide Sequences of 500 Mouse
 KIAA-Homologous cDNAs Identified by Screening of Terminal Sequences
 of cDNA Clones Randomly Sampled from Size-Fractionated Libraries
 DNA Res. 11, 205-218 (2004)

2 (bases 1 to 5551)

Okazaki,N., Kikuno,R.F., Nagase,T., Ohara,O. and Koga,H.

Direct Submission

Submitted (19-MAY-2004) Hisashi Koga, Kazusa DNA Research

Institute, Laboratory for Genome Informatics; 2-6-7

Kazusa-kamatari, Kisarazu, Chiba 292-0818, Japan

(E-mail:mouse@kazusa.or.jp, Tel:81-438-52-3919, Fax:81-438-52-3918)

The CREATE program supported by Japan science and technology

corporation; cDNA full insert sequencing; Kazusa DNA Research

Institute; cDNA library construction, clone selection and 5'- &

3'-end one pass sequencing.

Location/Qualifiers

FEATURES

source

1..5551

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="mfj04206"

/note="vector:modified pBC SK+"

1..5551

/gene="mKIAA1973"

<1..3097

/gene="mKIAA1973"

/note="CDS is predicted by in silico analysis. Start codon

is not identified."

/codon_start=2

/evidence=not experimental

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/protein_id="BAD32592.1"

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FSLLLCQEDWNITDFLLTENNPKFLESIIINITANLSSTKDLLSLFQVLENIRNST
PTMVFQCDMGSIROI FEMSTQFGLSPDLHWLVDGDSQNVLELRTGLPLGLIAHGKT
TQSFVEYVQDAMELAVARATMTIQPELALLPTMNCMDVKTLLTSQYLSRFLA
NTTFRGLSGSIKVGSTIVSSNFPFIWNLQYDPMKPMWTKTGLSQQVGGRI VMDSGIW
PQAOHKTTHFHPNKLHLRVVTLIEHPVFTREVDDEGLCPAGQLCDPMTNDSSIL
DSLSLSHSSNDTVPKFKKCCYGCIDLLLEQLAEDMNFDPOLYIVDGDCKYGAWKXGH
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GRTAATKPKCKWTFRLMNLWAI FCMFCLSTYTANLAAVMGEKIYEELSGIHDPKLIH
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SUFFRNSMGFQQLMVMYNTSNLSDNQRYIFNDEGQNLGTQTHQDIPLPFRRLPL
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ORIGIN

Query Match 72.6%; Score 19.6; DB 10; Length 5551;
 Best Local Similarity 84.6%; Pred. No. 79;
 Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGTACCCCTAGGCTTAGGC 27

Db 4186 TAGCCTAGTTCCTCCCTTTGTTAGGC 4211

RESULT 14

AL732521/c

LOCUS

DEFINITION

Mouse DNA sequence from clone RP23-134A17 on chromosome 4, complete

sequence.

ACCESSION AL732521

VERSION AL732521.18 GI:32879627

KEYWORDS HTG.

SOURCE Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

REFERENCE 1 (bases 1 to 207223)

AUTHORS Wallis,J.

Direct Submission

Submitted (16-JUL-2003) Wellcome Trust Sanger Institute, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquiries:

humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk

On Jul 16, 2003 this sequence version replaced gi:32480449.

Sequence from the Mouse Genome Sequencing Consortium whole genome

shotgun may have been used to confirm this sequence. Sequence data

from the whole genome shotgun alone has only been used where it has

a phred quality of at least 30.

----- Genome Center

Center: Wellcome Trust Sanger Institute

Center code: SC

Web site: http://www.sanger.ac.uk

Contact: humquery@sanger.ac.uk

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all

regions were either double-stranded or sequenced with an alternate

chemistry or covered by high quality data (i.e., phred quality >=

30); an attempt was made to resolve all sequencing problems, such

as compressions and repeats; all regions were covered by at least

one plasmid subclone or more than one M13 subclone; and the

assembly was confirmed by restriction digest, except on the rare

occasion of the clone being a YAC.
The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases:
Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep RP23-134A17 is from the RPCI-23 Mouse BAC Library constructed by the group of Pieter de Jong.
For further details see <http://www.chori.org/bacpac/home.htm>
VECTOR: pBACE3.6

FEATURES

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ORIGIN

Query Match 72.6%; Score 19.6; DB 10; Length 207223;
Best Local Similarity 84.6%; Pred. NO. 61;
Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
QY 2 TAGCCTAGCTACCCCTAGGCTTAGGC 27
|||||
Db 117297 TAGCCTAGCTTCCCTTTGTTAGGC 117272

RESULT 15

AC126959

LOCUS

AC126959 212481 bp DNA linear ROD 12-AUG-2004
Rattus norvegicus 8 BAC CH230-10P12 (Children's Hospital Oakland Research Institute) complete sequence.

AC126959

VERSION

AC126959.6 GI:49170123

KEYWORDS

Rattus norvegicus (Norway rat)

SOURCE

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus

1 (bases 1 to 212481)

Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,

Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,

Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,

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Watson, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J.,
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Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D. R.,
Holt, R. A., Smith, H. O., Weinstock, G. and Gibbs, R. A.

TITLE
JOURNAL

REFERENCE
2 (bases 1 to 212481)

AUTHORS
Worley, K.C.

TITLE
JOURNAL

Submitted (11-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 212481)

REFERENCE
AUTHORS

CONSTRM
JOURNAL

TITLE
JOURNAL

Submitted (10-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 212481)

REFERENCE
AUTHORS

TITLE
JOURNAL

Submitted (24-JUN-2004) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

REFERENCE
AUTHORS

TITLE
JOURNAL

Submitted (10-MAY-2003) Human Genome Sequencing Center, Department
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4 (bases 1 to 212481)

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Query Match 72.6%; Score 19.6; DB 10; Length 212481;
 Best Local Similarity 84.6%; Pred. No. 61;
 Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 TAGCCTAGTACCCCTAGGCTAGGC 27
 |||||
 DB 91228 TATTCTATCTATCCCTAGGCTAGGC 91253

Search completed: October 13, 2005, 19:37:32
 Job time : 1488 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 13, 2005, 17:44:53 ; Search time 1122 Seconds
(without alignments)
142.454 Million cell updates/sec

Title: US-09-744-097A-76

Perfect score: 27
Sequence: 1 gtgcctagctaccctagctctagcgc 27

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_16Dec04:
1: geneseqn1980s:
2: geneseqn1990s:
3: geneseqn2000s:
4: geneseqn2001as:
5: geneseqn2001bs:
6: geneseqn2002as:
7: geneseqn2002bs:
8: geneseqn2003as:
9: geneseqn2003bs:
10: geneseqn2003cs:
11: geneseqn2003ds:
12: geneseqn2004as:
13: geneseqn2004bs:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	27	100.0	27	3 AAA32245	Aaa32245 Green ter
2	27	100.0	27	3 AAA32253	Aaa32253 Distal sp
3	27	100.0	27	3 AAA32261	Aaa32261 Double di
4	27	100.0	27	3 AAA32262	Aaa32262 Double di
5	27	100.0	27	3 AAA32274	Aaa32274 One to tw
6	27	100.0	27	3 AAA32293	Aaa32293 Green set
7	27	100.0	27	3 AAA32255	Aaa32255 Proximal
8	27	100.0	27	3 AAA32243	Aaa32243 Distal sp
9	27	100.0	27	3 AAA32264	Aaa32264 Proximal
10	27	100.0	27	3 AAA32241	Aaa32241 Proximal
11	27	100.0	27	3 AAA32275	Aaa32275 One to tw
12	27	100.0	27	4 AAS14192	Aas14192 GenetAG F
13	27	100.0	27	4 AAS14194	Aas14194 First-GRE
14	27	100.0	27	4 AAS14183	Aas14183 Fragment
15	27	100.0	27	4 AAS14198	Aas14198 First-GRE
16	27	100.0	27	4 AAS14191	Aas14191 First-GRE
17	27	100.0	27	4 AAS14181	Aas14181 Fragment
18	27	100.0	29	4 AAS14197	Aas14197 First-GRE
19	27	100.0	31	4 AAS14190	Aas14190 First-GRE
20	27	100.0	36	4 AAS14195	Aas14195 First-GRE

C 21	27	100.0	42	3 AAA32311	Aaa32311 Anti-sens
C 22	27	100.0	42	4 AAS14178	Aas14178 Modified
C 23	27	100.0	57	3 AAA32233	Aaa32233 Green rep
C 24	27	100.0	72	3 AAA32310	Aaa32310 Anti-sens
C 25	25	92.6	25	3 AAA32225	Aaa32225 Terminato
C 26	25	92.6	25	3 AAA32188	Aaa32188 Terminato
C 27	25	92.6	25	3 AAA32177	Aaa32177 Distal li
C 28	25	92.6	37	3 AAA32184	Aaa32184 Overlap l
C 29	25	92.6	37	3 AAA32174	Aaa32174 Overlap o
C 30	25	92.6	49	3 AAA32220	Aaa32220 Reporter
C 31	25	92.6	57	3 AAA32222	Aaa32222 Proximal
C 32	25	92.6	57	3 AAA32223	Aaa32223 Distal su
C 33	25	92.6	57	3 AAA32186	Aaa32186 Distal su
C 34	25	92.6	57	3 AAA32185	Aaa32185 Proximal
C 35	25	92.6	63	3 AAA32173	Aaa32173 Target sp
C 36	25	92.6	63	3 AAA32183	Aaa32183 Target ol
C 37	24	88.9	24	3 AAA32175	Aaa32175 Proximal
C 38	24	88.9	24	3 AAA32182	Aaa32182 Terminato
C 39	24	88.9	24	3 AAA32179	Aaa32179 Proximal
C 40	24	88.9	40	3 AAA32199	Aaa32199 Second re
C 41	24	88.9	40	3 AAA32191	Aaa32191 Forward r
C 42	24	88.9	40	3 AAA32193	Aaa32193 Reverse r
C 43	24	88.9	40	3 AAA32204	Aaa32204 Forward r
C 44	24	88.9	40	3 AAA32197	Aaa32197 Reporter
C 45	24	88.9	40	3 AAA32206	Aaa32206 Reverse r

ALIGNMENTS

RESULT 1
AAA32245
ID AAA32245 standard; DNA; 27 BP.
XX
AC AAA32245;
XX 14-JUL-2000 (first entry)
XX Green terminator sequence used in gene-tag reporter construction.
DE Gene-tag reporter; detection; gene mapping; mutation identification;
KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX Synthetic.
XX WO200004192-A1.
XX 27-JAN-2000.
XX 16-JUL-1999; 99WO-US016242.
XX 17-JUL-1998; 98US-0093219P.
XX (UYEM-) UNIV EMORY.
XX Shafer DA;
XX WPI; 2000-182448/16.
XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for mapping genes or mutational analysis.
XX Example 12; Page 62; 164pp; English.
XX This sequence is used in the construction of the gene-tag reporter of the invention. The invention relates to a gene-tag reporter for joining a probe, alone or with a second gene-tag reporter. The gene-tag reporter comprises a labelled double-stranded polynucleotide sequence having one or more linkers that comprise a single stranded sequence hybridisable to a complement but not to the target probe. Also included in the invention is a reporter array, comprising at least two gene-tag reporters linked end to end by hybridisable linkers. The reporters are used in various new methods for detecting and mapping genes; identifying mutations and

CC variant nucleic acids, e.g. detecting rare mutations such as those in
 CC cancer cells or mutant viruses, but more generally in human diagnostics,
 CC forensics, genetic analysis, analysis of environmental samples or foods.
 CC The gene-tag reporters and associated probes, have a modular structure,
 CC allowing simple and inexpensive probe design, and are able to generate a
 CC distinctive signal, based on the mix and/or proportions of different
 CC signal components. Many targets can be analysed simultaneously, using
 CC many probes. Arrays of gene-tag reporters will provide signal
 CC amplification
 XX
 SQ Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 27; DB 3; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27
 Db 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27

RESULT 2
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 ID AAA32253 standard; DNA; 27 BP.
 XX
 AC AAA32253;
 XX
 DT 14-JUL-2000 (first entry)
 XX
 DE Distal spacer oligomer A used in gene-tag reporter construction.
 XX
 KW Gene-tag reporter; detection; gene mapping; mutation identification;
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
 XX
 OS Synthetic.
 XX
 PN WO200004192-A1.
 XX
 PD 27-JAN-2000.
 XX
 PF 16-JUL-1999; 99WO-US016242.
 XX
 PR 17-JUL-1998; 98US-0093219P.
 XX
 PA (UYEM-) UNIV EMORY.
 XX
 PI Shafer DA;
 XX
 DR WPI; 2000-182448/16.
 XX
 PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
 PT mapping genes or mutational analysis.
 XX
 PS Example 13; Page 67; 164pp; English.
 XX

CC This sequence is used in the construction of the gene-tag reporter of the
 CC invention. The invention relates to a gene-tag reporter for joining a
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
 CC comprises a labelled double-stranded polynucleotide sequence having one
 CC or more linkers that comprise a single stranded sequence hybridisable to
 CC a complement but not to the target probe. Also included in the invention
 CC is a reporter array, comprising at least two gene-tag reporters linked
 CC end to end by hybridisable linkers. The reporters are used in various new
 CC methods for detecting and mapping genes; identifying mutations and
 CC variant nucleic acids, e.g. detecting rare mutations such as those in
 CC cancer cells or mutant viruses, but more generally in human diagnostics,
 CC forensics, genetic analysis, analysis of environmental samples or foods.
 CC The gene-tag reporters and associated probes, have a modular structure,
 CC allowing simple and inexpensive probe design, and are able to generate a
 CC distinctive signal, based on the mix and/or proportions of different
 CC signal components. Many targets can be analysed simultaneously, using
 CC many probes. Arrays of gene-tag reporters will provide signal
 CC amplification

XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
 SQ
 Query Match 100.0%; Score 27; DB 3; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27
 Db 27 GTAGCCTAGTACCCCTAGGTCCTAGGC 1

RESULT 3
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 ID AAA32261 standard; DNA; 27 BP.
 XX
 AC AAA32261;
 XX
 DT 14-JUL-2000 (first entry)
 XX
 DE Double distal spacer oligomer A used in gene-tag reporter construction.
 XX
 KW Gene-tag reporter; detection; gene mapping; mutation identification;
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
 XX
 OS Synthetic.
 XX
 PN WO200004192-A1.
 XX
 PD 27-JAN-2000.
 XX
 PF 16-JUL-1999; 99WO-US016242.
 XX
 PR 17-JUL-1998; 98US-0093219P.
 XX
 PA (UYEM-) UNIV EMORY.
 XX
 PI Shafer DA;
 XX
 DR WPI; 2000-182448/16.
 XX
 PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
 PT mapping genes or mutational analysis.
 XX
 PS Example 14; Page 69; 164pp; English.
 XX

CC This sequence is used in the construction of the gene-tag reporter of the
 CC invention. The invention relates to a gene-tag reporter for joining a
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
 CC comprises a labelled double-stranded polynucleotide sequence having one
 CC or more linkers that comprise a single stranded sequence hybridisable to
 CC a complement but not to the target probe. Also included in the invention
 CC is a reporter array, comprising at least two gene-tag reporters linked
 CC end to end by hybridisable linkers. The reporters are used in various new
 CC methods for detecting and mapping genes; identifying mutations and
 CC variant nucleic acids, e.g. detecting rare mutations such as those in
 CC cancer cells or mutant viruses, but more generally in human diagnostics,
 CC forensics, genetic analysis, analysis of environmental samples or foods.
 CC The gene-tag reporters and associated probes, have a modular structure,
 CC allowing simple and inexpensive probe design, and are able to generate a
 CC distinctive signal, based on the mix and/or proportions of different
 CC signal components. Many targets can be analysed simultaneously, using
 CC many probes. Arrays of gene-tag reporters will provide signal
 CC amplification

XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
 SQ
 Query Match 100.0%; Score 27; DB 3; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 27
 Db 1 GTAGCCTAGTACCCCTAGGTCCTAGGC 1

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Db      27 GTAGCCTAGCTACCCCTAGGCTAGGC 1
RESULT 4
AAA32262/c
ID      AAA32262 standard; DNA; 27 BP.
XX
AC      AAA32262;
XX
XX      14-JUL-2000 (first entry)
DT      14-JUL-2000 (first entry)
DE      One to two multilinker A used in gene-tag reporter construction.
XX
XX      Gene-tag reporter; detection; gene mapping; mutation identification;
KW      cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
OS      Synthetic.
XX
PN      WO200004192-A1.
XX
PD      27-JAN-2000.
XX
PF      16-JUL-1999; 99WO-US016242.
XX
PR      17-JUL-1998; 98US-0093219P.
XX
PA      (UYEM-) UNIV EMORY.
XX
PI      Shafer DA;
XX
XX      WPI; 2000-182448/16.
XX
XX      New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT      mapping genes or mutational analysis.
XX
PS      Example 15; Page 70; 164pp; English.
XX
CC      This sequence is used in the construction of the gene-tag reporter of the
CC      invention. The invention relates to a gene-tag reporter for joining a
CC      probe, alone or with a second gene-tag reporter. The gene-tag reporter
CC      comprises a labelled double-stranded polynucleotide sequence having one
CC      or more linkers that comprise a single stranded sequence hybridisable to
CC      a complement but not to the target probe. Also included in the invention
CC      is a reporter array, comprising at least two gene-tag reporters linked
CC      end to end by hybridisable linkers. The reporters are used in various new
CC      methods for detecting and mapping genes; identifying mutations and
CC      variant nucleic acids, e.g. detecting rare mutations such as those in
CC      cancer cells or mutant viruses, but more generally in human diagnostics,
CC      forensics, genetic analysis, analysis of environmental samples or foods.
CC      The gene-tag reporters and associated probes, have a modular structure,
CC      allowing simple and inexpensive probe design, and are able to generate a
CC      distinctive signal, based on the mix and/or proportions of different
CC      signal components. Many targets can be analysed simultaneously, using
CC      many probes. Arrays of gene-tag reporters will provide signal
CC      amplification
XX
SQ      Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
        Query Match      100.0%; Score 27; DB 3; Length 27;
        Best Local Similarity 100.0%; Pred. No. 0.0075;
        Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTAGCCTAGCTACCCCTAGGCTAGGC 27
        |||||
Db      27 GTAGCCTAGCTACCCCTAGGCTAGGC 1

RESULT 6
AAA32293/c
ID      AAA32293 standard; DNA; 27 BP.
XX
AC      AAA32293;
XX
XX      14-JUL-2000 (first entry)
DT
DE
XX      Green set first linker A used in gene-tag reporter construction.
XX
XX      Gene-tag reporter; detection; gene mapping; mutation identification;
KW      cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
OS      Synthetic.
XX

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PN WO200004192-A1.
PD 27-JAN-2000.
XX 16-JUL-1999; 99WO-US016242.
XX 17-JUL-1998; 98US-0093219P.
XX (UYEM-) UNIV EMORY.
PA Shafer DA;
PI WPI; 2000-182448/16.
DR New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT mapping genes or mutational analysis.
XX Example 16; Page 72; 164pp; English.
XX This sequence is used in the construction of the gene-tag reporter of the
CC invention. The invention relates to a gene-tag reporter for joining a
CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
CC comprises a labelled double-stranded polynucleotide sequence having one
CC or more linkers that comprise a single stranded sequence hybridisable to
CC a complement but not to the target probe. Also included in the invention
CC is a reporter array, comprising at least two gene-tag reporters linked
CC end to end by hybridisable linkers. The reporters are used in various new
CC methods for detecting and mapping genes; identifying mutations and
CC variant nucleic acids, e.g. detecting rare mutations such as those in
CC cancer cells or mutant viruses, but more generally in human diagnostics,
CC forensics, genetic analysis, analysis of environmental samples or foods.
CC The gene-tag reporters and associated probes, have a modular structure,
CC allowing simple and inexpensive probe design, and are able to generate a
CC distinctive signal, based on the mix and/or proportions of different
CC signal components. Many targets can be analysed simultaneously, using
CC many probes. Arrays of gene-tag reporters will provide signal
CC amplification
XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
SQ Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
Db |||||
27 GTAGCCTAGCTACCCCTAGTCTAGGC 1

RESULT 7
AAA32255
ID AAA32255 standard; DNA; 27 BP.
XX AAA32255;
AC 14-JUL-2000 (first entry)
XX Proximal spacer oligomer A used in gene-tag reporter construction.
DE Gene-tag reporter; detection; gene mapping; mutation identification;
KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX Synthetic.
OS WO200004192-A1.
XX 27-JAN-2000.
XX 16-JUL-1999; 99WO-US016242.
XX 17-JUL-1998; 98US-0093219P.
XX (UYEM-) UNIV EMORY.
PA Shafer DA;
PI WPI; 2000-182448/16.
DR New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT mapping genes or mutational analysis.
XX Example 16; Page 72; 164pp; English.
XX This sequence is used in the construction of the gene-tag reporter of the
CC invention. The invention relates to a gene-tag reporter for joining a
CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
CC comprises a labelled double-stranded polynucleotide sequence having one
CC or more linkers that comprise a single stranded sequence hybridisable to
CC a complement but not to the target probe. Also included in the invention
CC is a reporter array, comprising at least two gene-tag reporters linked
CC end to end by hybridisable linkers. The reporters are used in various new
CC methods for detecting and mapping genes; identifying mutations and
CC variant nucleic acids, e.g. detecting rare mutations such as those in
CC cancer cells or mutant viruses, but more generally in human diagnostics,
CC forensics, genetic analysis, analysis of environmental samples or foods.
CC The gene-tag reporters and associated probes, have a modular structure,
CC allowing simple and inexpensive probe design, and are able to generate a
CC distinctive signal, based on the mix and/or proportions of different
CC signal components. Many targets can be analysed simultaneously, using
CC many probes. Arrays of gene-tag reporters will provide signal
CC amplification
XX Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
SQ Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
Db |||||
27 GTAGCCTAGCTACCCCTAGTCTAGGC 1

RESULT 8
AAA32243/C
ID AAA32243 standard; DNA; 27 BP.
XX AAA32243;
AC 14-JUL-2000 (first entry)
XX Distal spacer oligomer A used in gene-tag reporter construction.
DE Gene-tag reporter; detection; gene mapping; mutation identification;
KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX Synthetic.
OS WO200004192-A1.
XX 27-JAN-2000.
XX 16-JUL-1999; 99WO-US016242.
XX 17-JUL-1998; 98US-0093219P.
XX (UYEM-) UNIV EMORY.
PA Shafer DA;
PI WPI; 2000-182448/16.
DR New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT mapping genes or mutational analysis.
XX Example 12; Page 62; 164pp; English.

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XX Shafer DA;
PI WPI; 2000-182448/16.
DR New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT mapping genes or mutational analysis.
XX Example 13; Page 67; 164pp; English.
XX This sequence is used in the construction of the gene-tag reporter of the
CC invention. The invention relates to a gene-tag reporter for joining a
CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
CC comprises a labelled double-stranded polynucleotide sequence having one
CC or more linkers that comprise a single stranded sequence hybridisable to
CC a complement but not to the target probe. Also included in the invention
CC is a reporter array, comprising at least two gene-tag reporters linked
CC end to end by hybridisable linkers. The reporters are used in various new
CC methods for detecting and mapping genes; identifying mutations and
CC variant nucleic acids, e.g. detecting rare mutations such as those in
CC cancer cells or mutant viruses, but more generally in human diagnostics,
CC forensics, genetic analysis, analysis of environmental samples or foods.
CC The gene-tag reporters and associated probes, have a modular structure,
CC allowing simple and inexpensive probe design, and are able to generate a
CC distinctive signal, based on the mix and/or proportions of different
CC signal components. Many targets can be analysed simultaneously, using
CC many probes. Arrays of gene-tag reporters will provide signal
CC amplification
XX Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
SQ Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
Db |||||
1 GTAGCCTAGCTACCCCTAGTCTAGGC 27

RESULT 8
AAA32243/C
ID AAA32243 standard; DNA; 27 BP.
XX AAA32243;
AC 14-JUL-2000 (first entry)
XX Distal spacer oligomer A used in gene-tag reporter construction.
DE Gene-tag reporter; detection; gene mapping; mutation identification;
KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX Synthetic.
OS WO200004192-A1.
XX 27-JAN-2000.
XX 16-JUL-1999; 99WO-US016242.
XX 17-JUL-1998; 98US-0093219P.
XX (UYEM-) UNIV EMORY.
PA Shafer DA;
PI WPI; 2000-182448/16.
DR New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT mapping genes or mutational analysis.
XX Example 12; Page 62; 164pp; English.

```

xx This sequence is used in the construction of the gene-tag reporter of the
 CC invention. The invention relates to a gene-tag reporter for joining a
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
 CC comprises a labelled double-stranded polynucleotide sequence having one
 CC or more linkers that comprise a single stranded sequence hybridisable to
 CC a complement but not to the target probe. Also included in the invention
 CC is a reporter array, comprising at least two gene-tag reporters linked
 CC end to end by hybridisable linkers. The reporters are used in various new
 CC methods for detecting and mapping genes; identifying mutations and
 CC variant nucleic acids, e.g. detecting rare mutations such as those in
 CC cancer cells or mutant viruses, but more generally in human diagnostics,
 CC forensics, genetic analysis, analysis of environmental samples or foods.
 CC The gene-tag reporters and associated probes, have a modular structure,
 CC allowing simple and inexpensive probe design, and are able to generate a
 CC distinctive signal, based on the mix and/or proportions of different
 CC signal components. Many targets can be analysed simultaneously, using
 CC many probes. Arrays of gene-tag reporters will provide signal
 CC amplification

SQ Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
 Query Match 100.0%; Score 27; DB 3; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGGCTTAGGC 27
 DB 27 GTAGCCTAGCTACCCCTAGGCTTAGGC 1

RESULT 9
 AAA32264
 ID AAA32264 standard; DNA; 27 BP.
 AC AAA32264;
 XX
 XX 14-JUL-2000 (first entry)
 DT
 XX Proximal spacer oligomer A used in gene-tag reporter construction.
 DE
 XX Gene-tag reporter; detection; gene mapping; mutation identification;
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
 XX Synthetic.
 OS
 XX WO200004192-A1.
 FN
 XX 27-JAN-2000.
 PD
 XX 16-JUL-1999; 99WO-US016242.
 PF
 XX 17-JUL-1998; 98US-0093219P.
 PR
 XX (UYEM-) UNIV EMORY.
 PA
 XX Shafer DA;
 PI
 XX WPI; 2000-182448/16.
 DR
 XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
 PT mapping genes or mutational analysis.
 PT
 XX Example 14; Page 69; 164pp; English.
 PS
 XX This sequence is used in the construction of the gene-tag reporter of the
 CC invention. The invention relates to a gene-tag reporter for joining a
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
 CC comprises a labelled double-stranded polynucleotide sequence having one
 CC or more linkers that comprise a single stranded sequence hybridisable to
 CC a complement but not to the target probe. Also included in the invention
 CC is a reporter array, comprising at least two gene-tag reporters linked
 CC end to end by hybridisable linkers. The reporters are used in various new

CC methods for detecting and mapping genes; identifying mutations and
 CC variant nucleic acids, e.g. detecting rare mutations such as those in
 CC cancer cells or mutant viruses, but more generally in human diagnostics,
 CC forensics, genetic analysis, analysis of environmental samples or foods.
 CC The gene-tag reporters and associated probes, have a modular structure,
 CC allowing simple and inexpensive probe design, and are able to generate a
 CC distinctive signal, based on the mix and/or proportions of different
 CC signal components. Many targets can be analysed simultaneously, using
 CC many probes. Arrays of gene-tag reporters will provide signal
 CC amplification

SQ Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
 Query Match 100.0%; Score 27; DB 3; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGCTACCCCTAGGCTTAGGC 27
 DB 1 GTAGCCTAGCTACCCCTAGGCTTAGGC 27

RESULT 10
 AAA32241
 ID AAA32241 standard; DNA; 27 BP.
 AC AAA32241;
 XX
 XX 14-JUL-2000 (first entry)
 DT
 XX Proximal spacer oligomer A used in gene-tag reporter construction.
 DE
 XX Gene-tag reporter; detection; gene mapping; mutation identification;
 KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
 XX Synthetic.
 OS
 XX WO200004192-A1.
 FN
 XX 27-JAN-2000.
 PD
 XX 16-JUL-1999; 99WO-US016242.
 PF
 XX 17-JUL-1998; 98US-0093219P.
 PR
 XX (UYEM-) UNIV EMORY.
 PA
 XX Shafer DA;
 PI
 XX WPI; 2000-182448/16.
 DR
 XX New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
 PT mapping genes or mutational analysis.
 PT
 XX Example 12; Page 62; 164pp; English.
 PS
 XX This sequence is used in the construction of the gene-tag reporter of the
 CC invention. The invention relates to a gene-tag reporter for joining a
 CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
 CC comprises a labelled double-stranded polynucleotide sequence having one
 CC or more linkers that comprise a single stranded sequence hybridisable to
 CC a complement but not to the target probe. Also included in the invention
 CC is a reporter array, comprising at least two gene-tag reporters linked
 CC end to end by hybridisable linkers. The reporters are used in various new
 CC methods for detecting and mapping genes; identifying mutations and
 CC variant nucleic acids, e.g. detecting rare mutations such as those in
 CC cancer cells or mutant viruses, but more generally in human diagnostics,
 CC forensics, genetic analysis, analysis of environmental samples or foods.
 CC The gene-tag reporters and associated probes, have a modular structure,
 CC allowing simple and inexpensive probe design, and are able to generate a
 CC distinctive signal, based on the mix and/or proportions of different
 CC signal components. Many targets can be analysed simultaneously, using
 CC many probes. Arrays of gene-tag reporters will provide signal

CC amplification
XX Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
SQ Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27
|||
Db 1 GTAGCCTAGTACCCCTAGTCTAGGC 27

RESULT 11
AAA32275/c
ID AAA32275 standard; DNA; 27 BP.
XX
AC AAA32275;
XX
DT 14-JUL-2000 (first entry)
XX
DE One to two multilinker B used in gene-tag reporter construction.
XX
KW Gene-tag reporter; detection; gene mapping; mutation identification;
KW cancer; mutant virus; human diagnostic; forensic; genetic analysis; ss.
XX
OS Synthetic.
XX WO200004192-A1.
XX
PD 27-JAN-2000.
XX
PF 16-JUL-1999; 99WO-US016242.
XX
PR 17-JUL-1998; 98US-0093219P.
XX
PA (UYEM-) UNIV EMORY.
XX
PI Shafer DA;
XX
WPI; 2000-182448/16.
XX
PT New Gene-tag reporter for joining to nucleic acid probe, used, e.g. for
PT mapping genes or mutational analysis.
XX
PS Example 15; Page 70; 164pp; English.
XX
CC This sequence is used in the construction of the gene-tag reporter of the
CC invention. The invention relates to a gene-tag reporter for joining a
CC probe, alone or with a second gene-tag reporter. The gene-tag reporter
CC comprises a labelled double-stranded polynucleotide sequence having one
CC or more linkers that comprise a single stranded sequence hybridisable to
CC a complement but not to the target probe. Also included in the invention
CC is a reporter array, comprising at least two gene-tag reporters linked
CC end to end by hybridisable linkers. The reporters are used in various new
CC methods for detecting and mapping genes; identifying mutations and
CC variant nucleic acids, e.g. detecting rare mutations such as those in
CC cancer cells or mutant viruses, but more generally in human diagnostics,
CC forensics, genetic analysis, analysis of environmental samples or foods.
CC The gene-tag reporters and associated probes, have a modular structure,
CC allowing simple and inexpensive probe design, and are able to generate a
CC distinctive signal, based on the mix and/or proportions of different
CC signal components. Many targets can be analysed simultaneously, using
CC many probes. Arrays of gene-tag reporters will provide signal
CC amplification
XX
SQ Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 27; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27
|||
Db 1 GTAGCCTAGTACCCCTAGTCTAGGC 27

RESULT 13
AAA14194/c
ID AAA14194 standard; DNA; 27 BP.
XX
AC AAA14192;
XX
DT 18-DEC-2001 (first entry)
XX
DE GeneTAG First-GREEN primer used in construction of probe sets.
XX
KW WRAP-Probe; gene expression array; global amplification; RNA array; ss;
KW tissue microarray; drug discovery assay; reporter binding site; forensic;
KW diagnostic; genomic analysis; universal linker; PCR primer.
XX
OS Synthetic.
XX WO200166802-A1.
XX
PD 13-SEP-2001.
XX
PF 09-MAR-2001; 2001WO-US007508.
XX
PR 09-MAR-2000; 2000US-0187982P.
XX
PA (GENE-) GENETAG TECHNOLOGY INC.
XX
PI Shafer DA;
XX
WPI; 2001-596845/67.
XX
PT Novel probe sets with common universal linkers at one or both ends (WRAP
PT probes) for gene expression arrays to provide global amplification of
PT probe set and to provide common equivalent signaling regardless of
PT length.
XX
PS Example 3; Page 59; 97pp; English.
XX
CC The invention relates to a probe set for gene expression arrays to
CC provide common equivalent signalling per probe and global amplification
CC of the set. The probe set has a pool of modified cDNA probes, each probe
CC having a central target specific segment copied from a portion of a
CC single mRNA transcript and a universal linker (a WRAP-probe) located on
CC one or both terminal ends. The universal linker has reporter binding
CC sites to join common reporters to the probes and primer binding sites to
CC copy and amplify the probe. The probes and reporters are useful in
CC diagnostic or drug discovery assays for a wide range of biomedical
CC samples, including detection of nucleic acids and gene expression
CC profiles in human diagnostics, forensics and genomic analysis. The
CC methods are useful for amplifying and identifying any unknown DNA
CC fragment and also for improving sensitivity with tissue microarrays or
CC RNA arrays. The methods improve the quantification of gene expression and
CC allow highly improved detection of rare transcripts or very small
CC samples. This sequence represents a GenetAG First-GREEN primer used in
CC the construction of probe sets
XX
SQ Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 27; DB 4; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGTCTAGGC 27
|||
Db 27 GTAGCCTAGTACCCCTAGTCTAGGC 1

RESULT 13
AAA14194/c
ID AAA14194 standard; DNA; 27 BP.

xx AAS14194;
 xx 18-DEC-2001 (first entry)
 xx First-GREEN Chiptag primer used in construction of probe sets.
 DE
 xx WRAP-Probe; gene expression array; global amplification; RNA array; ss;
 xx tissue microarray; drug discovery assay; reporter binding site; forensic;
 xx diagnostic; genomic analysis; universal linker; PCR primer.
 xx Synthetic.
 OS
 xx WO200166802-A1.
 xx 13-SEP-2001.
 xx 09-MAR-2001; 2001WO-US007508.
 xx 09-MAR-2000; 2000US-0187982P.
 xx (GENE-) GENETAG TECHNOLOGY INC.
 xx Shafer DA;
 xx WPI; 2001-596845/67.
 xx Novel probe sets with common universal linkers at one or both ends (WRAP
 PT probes) for gene expression arrays to provide global amplification of
 PT probe set and to provide common equivalent signaling regardless of
 PT length.
 xx Example 4; Page 60; 97pp; English.
 xx The invention relates to a probe set for gene expression arrays to
 CC provide common equivalent signaling per probe and global amplification
 CC of the set. The probe set has a pool of modified cDNA probes, each probe
 CC having a central target specific segment copied from a portion of a
 CC single mRNA transcript and a universal linker (a WRAP-Probe) located on
 CC one or both terminal ends. The universal linker has reporter binding
 CC sites to join common reporters to the probes and primer binding sites to
 CC copy and amplify the probe. The probes and reporters are useful in
 CC diagnostic or drug discovery assays for a wide range of biomedical
 CC samples, including detection of nucleic acids and gene expression
 CC profiles in human diagnostics, forensics and genomic analysis. The
 CC methods are useful for amplifying and identifying any unknown DNA
 CC fragment and also for improving sensitivity with tissue microarrays or
 CC RNA arrays. The methods improve the quantification of gene expression and
 CC allow highly improved detection of rare transcripts or very small
 CC samples. This sequence represents a First-GREEN Chiptag primer used in
 CC the construction of probe sets
 xx
 SQ Sequence 27 BP; 6 A; 7 C; 9 G; 5 T; 0 U; 0 Other;
 Query Match 100.0%; Score 27; DB 4; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
 Db 27 GTAGCCTAGCTACCCCTAGTCTAGGC 1
 RESULT 14
 AAS14183
 ID AAS14183 standard; DNA; 27 BP.
 xx AAS14183;
 xx 18-DEC-2001 (first entry)
 xx Fragment #1 of PCR primer GR-SPC-F used in construction of probe sets.
 xx

KW WRAP-Probe; gene expression array; global amplification; RNA array; ss;
 KW tissue microarray; drug discovery assay; reporter binding site; forensic;
 KW diagnostic; genomic analysis; universal linker; PCR primer.
 XX Synthetic.
 OS
 xx WO200166802-A1.
 xx 13-SEP-2001.
 xx 09-MAR-2001; 2001WO-US007508.
 xx 09-MAR-2000; 2000US-0187982P.
 xx (GENE-) GENETAG TECHNOLOGY INC.
 xx Shafer DA;
 xx WPI; 2001-596845/67.
 xx Novel probe sets with common universal linkers at one or both ends (WRAP
 PT probes) for gene expression arrays to provide global amplification of
 PT probe set and to provide common equivalent signaling regardless of
 PT length.
 xx Disclosure; Page 90; 97pp; English.
 xx The invention relates to a probe set for gene expression arrays to
 CC provide common equivalent signaling per probe and global amplification
 CC of the set. The probe set has a pool of modified cDNA probes, each probe
 CC having a central target specific segment copied from a portion of a
 CC single mRNA transcript and a universal linker (a WRAP-Probe) located on
 CC one or both terminal ends. The universal linker has reporter binding
 CC sites to join common reporters to the probes and primer binding sites to
 CC copy and amplify the probe. The probes and reporters are useful in
 CC diagnostic or drug discovery assays for a wide range of biomedical
 CC samples, including detection of nucleic acids and gene expression
 CC profiles in human diagnostics, forensics and genomic analysis. The
 CC methods are useful for amplifying and identifying any unknown DNA
 CC fragment and also for improving sensitivity with tissue microarrays or
 CC RNA arrays. The methods improve the quantification of gene expression and
 CC allow highly improved detection of rare transcripts or very small
 CC samples. This sequence represents a fragment of a PCR primer used in the
 CC construction of probe sets
 xx
 SQ Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
 Query Match 100.0%; Score 27; DB 4; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.0075;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
 Db 1 GTAGCCTAGCTACCCCTAGTCTAGGC 27
 RESULT 15
 AAS14198
 ID AAS14198 standard; DNA; 27 BP.
 xx AAS14198;
 xx 18-DEC-2001 (first entry)
 xx First-GREEN random adapter part 2 used in construction of probe sets.
 xx WRAP-Probe; gene expression array; global amplification; RNA array; ss;
 KW tissue microarray; drug discovery assay; reporter binding site; forensic;
 KW diagnostic; genomic analysis; universal linker.
 XX Synthetic.
 OS
 xx WO200166802-A1.

XX 13-SEP-2001.
PD
XX
PF 09-MAR-2001; 2001WO-US007508.
XX
PR 09-MAR-2000; 2000US-0187982P.
XX
XX (GENE-) GENETAG TECHNOLOGY INC.
XX
XX Shafer DA;
PI
XX
DR WPI; 2001-596845/67.
XX
PT Novel probe sets with common universal linkers at one or both ends (WRAP
PT probes) for gene expression arrays to provide global amplification of
PT probe set and to provide common equivalent signaling regardless of
PT length.
XX
PS Example 7; Page 65; 97pp; English.
XX
CC The invention relates to a probe set for gene expression arrays to
CC provide common equivalent signalling per probe and global amplification
CC of the set. The probe set has a pool of modified cDNA probes, each probe
CC having a central target specific segment copied from a portion of a
CC single mRNA transcript and a universal linker (a WRAP-Probe) located on
CC one or both terminal ends. The universal linker has reporter binding
CC sites to join common reporters to the probes and primer binding sites to
CC copy and amplify the probe. The probes and reporters are useful in
CC diagnostic or drug discovery assays for a wide range of biomedical
CC samples, including detection of nucleic acids and gene expression
CC profiles in human diagnostics, forensics and genomic analysis. The
CC methods are useful for amplifying and identifying any unknown DNA
CC fragment and also for improving sensitivity with tissue microarrays or
CC RNA arrays. The methods improve the quantification of gene expression and
CC allow highly improved detection of rare transcripts or very small
CC samples. This sequence represents a random adapter fragment used in the
CC construction of probe sets
XX
SQ Sequence 27 BP; 5 A; 9 C; 7 G; 6 T; 0 U; 0 Other;
Query Match 100.0%; Score 27; DB 4; Length 27;
Best Local Similarity 100.0%; Pred No. 0.0075;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTAGCCTAGCTACCCCTAGGTCTAGGC 27
Db 1 GTAGCCTAGCTACCCCTAGGTCTAGGC 27

Search completed: October 13, 2005, 19:12:37
Job time : 1123 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 13, 2005, 18:25:43 ; Search time 7556 Seconds
(without alignments)
136.016 Million cell updates/sec

Title: US-09-744-097A-76

Perfect score: 27

Sequence: 1 gtagctagtagtacccttaggtctagc 27

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68473088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum-Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: gb_est1.*
2: gb_est2.*
3: gb_est3.*
4: gb_est4.*
5: gb_est5.*
6: gb_est6.*
7: gb_est7.*
8: gb_gss1.*
9: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19.6	72.6	297	2	BB274268
2	19.6	72.6	623	1	AV325957
3	19.6	72.6	801	2	BE394439
4	19.6	72.6	896	2	BE314562
5	19.6	72.6	3378	3	AK032394
6	19.2	71.1	115	9	CG805069
7	19.2	71.1	115	9	CG805158
8	19.2	71.1	419	8	A2636851
9	19.2	71.1	453	9	CG684396
10	19.2	71.1	630	8	B2631002
11	19.2	71.1	672	8	B2630997
12	19.2	71.1	683	8	B2633850
13	19.2	71.1	705	8	BH837552
14	19.2	71.1	725	8	B2633854
15	19.2	71.1	745	9	CG351990
16	19.2	71.1	747	9	CC626350
17	19.2	71.1	828	9	CC686399
18	19.2	71.1	839	9	CG306780
19	19.2	71.1	875	9	CG227499
20	19.2	71.1	902	9	CR222206
21	19.2	71.1	951	8	CC007567
22	19	70.4	478	7	CO781043
23	19	70.4	562	8	B2306625
24	19	70.4	774	8	CC122296

25	19	70.4	779	9	CC874542
26	19	70.4	786	9	CG063671
27	19	70.4	864	9	CG180375
28	19	70.4	865	9	CG180377
29	18.8	69.6	535	8	AZ500954
30	18.8	69.6	578	8	AZ492835
31	18.8	69.6	671	9	CR270282
32	18.8	69.6	675	9	CR273252
33	18.8	69.6	692	9	AG293534
34	18.8	69.6	711	9	AG286334
35	18.8	69.6	720	8	BZ257884
36	18.8	69.6	721	9	AG567157
37	18.6	68.9	806	7	CK316863
38	18.6	68.9	837	4	BG211849
39	18.6	68.9	869	8	BZ785030
40	18.6	68.9	880	8	BZ401569
41	18.6	68.9	954	8	CC295826
42	18.6	68.9	998	9	CG424886
43	18.6	68.9	1039	9	CC809843
44	18.2	67.4	406	8	AZ508009
45	18.2	67.4	428	6	CB097308

ALIGNMENTS

BB274268 297 bp mRNA linear EST 07-JUL-2000
BB274268 RIKEN full-length enriched, 10 days neonate cortex Mus
musculus cDNA clone AB30087F07 3', mRNA sequence.

BB274268

BB274268.1 GI:8971289

EST

Mus musculus (house mouse)

Mus musculus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

1 (bases 1 to 297)

REFERENCE

AUTHORS

Konno, H., Aizawa, K., Akahira, S., Akiyama, J., Arakawa, T.,
Carninci, P., Endo, T., Fukuda, S., Fukunishi, Y., Hara, A., Hayatsu, N.,
Hirozane, T., Hori, F., Ishii, Y., Ishikawa, J., Ishikawa, T., Itoh, M.,
Izawa, M., Kadota, K., Kagawa, I., Kai, C., Kawai, J., Kikuchi, N.,
Kiyosawa, H., Kojima, Y., Kondo, S., Koya, S., Kurihara, C.,
Kusakabe, M., Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H.,
Okazaki, Y., Ono, T., Owa, C., Saito, H., Sakai, C., Sato, K.,
Shibata, K., Shibata, Y., Shigemoto, Y., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Suganara, Y., Suzuki, H., Tagawa, A.,
Takahashi, F., Tomimaga, N., Toya, T., Tsunoda, Y., Watanabe, S.,
Watanabe, S., Yamamura, T., Yamana, I., Yano, R., Yasunishi, A.,
Yokota, T., Yoshida, K., Yoshiki, A., Yoshino, M., Muramatsu, M. and
Hayashizaki, Y.

RIKEN Mouse ESTs (Konno, H., et al.)

TITLE

JOURNAL

COMMENT

Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
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The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@sc.riken.jp, URL: http://genome.gsc.riken.jp/
Carninci, P., Nishiyama, Y., Westover, A., Itoh, M., Nagao, S.,
Sasaki, N., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Thermosensitization and thermoactivation of thermolabile enzymes by
trehalose and its application for the synthesis of full length
cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)
Itoh, M., Katsunari, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J.,
Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M.,
Okazaki, Y. and Hayashizaki, Y.
Automated filtration-based high-throughput plasmid preparation
system. Genome Res. 9 (5), 463-470 (1999)
Carninci, P. and Hayashizaki, Y.

High-efficiency full-length cDNA cloning. *Methods Enzymol.* 303, 19-44 (1999)
Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

FEATURES

```

1. .297
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="A830087F07"
/tissue_type="cortex"
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/lab_host="DH10B"
/clone_lib="RIKEN full-length enriched, 10 days neonate
cortex"
/notes="Site 1: SalI; Site 2: BamHI; cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN, Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer [5'
GAGAGAGAGAGATCCAGAGCTCTTTTTTTTTTTTNN 3'], cDNA was
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. cDNA went through one round of normalization
to Rot = 20.0 and subtraction to Rot = 459.0. Second
strand cDNA was prepared with the primer adaptor of
sequence [5' GAGAGAGAGATTCGAGTTAAATTAAATATATCCCCCCCCCCCC
3']. cDNA was cleaved with XhoI and BamHI. Vector: a
modified pBluescript KS(+) after bulk excision from Lambda
FLC I."

```

ORIGIN

```

atch          72.6%; Score 19.6; DB 2; Length 297;
al Similarity 84.6%; Pred. No. 2.9e+02;
22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

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RESULT 2

RESOLLI_2	AV325957	623 bp	mRNA	linear	EST 24-OCT-2001
LOCUS	AV325957	RIKEN full-length enriched,	adult male medulla oblongata		
DEFINITION	Mus musculus cDNA clone G330407C01 3', mRNA sequence.				
ACCESSION	AV325957				
VERSION	AV325957.2	GI:16395524			
KEYWORDS	EST.				

REFERENCE
AUTHORS

Hara, A., Hiramoto, K., Hori, F., Iehii, Y., Ito, M., Kawai, J.,
Kanno, H., Kouda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K.,
Onno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K.,
Sano, H., Sasaki, D., Shibata, K., Shingawa, A., Shiraki, I.,
Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,
Takeda, Y., Tanaka, T., Toyota, T., Muramatsu, M. and Hayashizaki, Y.
RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
Unpublished (2001)
On Nov 11, 1999 this sequence version replaced gi:6366009.

TITLE	JOURNAL	COMMENT
1. The Role of the Teacher in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 1, pp. 1-10
2. The Impact of Technology on Education	Journal of Educational Technology	1985, Vol. 10, No. 2, pp. 15-25
3. The Importance of Parental Involvement	Journal of Educational Psychology	1990, Vol. 82, No. 3, pp. 201-210
4. The Effect of Teacher Expectations on Student Achievement	Journal of Educational Research	1995, Vol. 98, No. 4, pp. 301-310
5. The Role of the School in the Community	Journal of Educational Research	2000, Vol. 103, No. 1, pp. 1-10

CONTACT: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
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Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gsc.riken.jp, URL: <http://genome.gsc.riken.jp/>
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Ito, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. *Genome Res.* 10 (10), 1617-1630 (2000).
wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T.,
Matsuda, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A.,
and Hayashizaki, Y.

and Hayashizaki, Y. RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)

Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y. Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)

Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamanaka, I., Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and Hayashizaki, Y. Computational Analysis of Full-Length Mouse cDNAs: Compared with Human Genome Sequences *Mamm. Genome*. 12, 673-677 (2001)

Please visit our web site (<http://genome.gsc.riken.go.jp/>) for further details.

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

FEATURES

source

```

1. .623
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/note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN, Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5' GAGAGAGAGGATCGACGAGCTCTTTTTCCTTTTTTN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 10.0 and subtraction to Rot = 100.0. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGATTCGAGTTAATAATTAAATCCCCCCCCCCC 3']. cDNA was cloned into the XhoI and BamHI sites. Vector: a modified phuscript KS(+) after bulk excision from Lambda PUC I. Cloning sites, 5' end: SalI; 3' end: BamHI."

```

ORIGIN

Query Match	72.6%	Score 19.6	DB 1	Length 623
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Matches	22	Conservative 0	Mismatches 4	Indels 0
2 TAGCCTAGCTACCCCTAGGCTAGGC 27				
569 TAGCCTAGCTTCCCTTTGTTTAGGC 594				

RESULT 3

MEDLINE
PUBMED
REFERENCE
4
AUTHORS
The RIKEN Genome Exploration Research Group Phase II Team and the
FANTOM Consortium.
TITLE
Functional annotation of a full-length mouse cDNA collection
JOURNAL
Nature 409, 685-690 (2001)
REFERENCE
5
AUTHORS
The FANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
JOURNAL
Nature 420, 563-573 (2002)
REFERENCE
6
AUTHORS
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T.,
Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Katoh, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M.,
Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,
Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N.,
Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A.,
Muramatsu, M., and Hayashizaki, Y.

TITLE
JOURNAL
Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp,
URL: http://genome.gsc.riken.jp/, Tel: 81-45-503-9222,
Fax: 81-45-503-9216)
COMMENT
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site for further details.
URL: http://genome.gsc.riken.jp/
URL: http://fantom.gsc.riken.jp/
FEATURES
Location/Qualifiers
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Matches 22; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 2 TAGCTAGCTACCCCTAGGCTAGGC 27
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Db 851 TAGCTAGCTTCCCTTTGTTAGGC 876
|||||

RESULT 6
CG805069 115 bp DNA linear GSS 10-NOV-2003
LOCUS
DEFINITION
1118057A09.x1 1118 - RescueMu Grid S Zea mays genomic, genomic
survey sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Zea mays

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Walbot, V.
Unpublished (2001)
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site so sequence was trimmed. Post-ligation
sequence submitted separately.
Plate: 1118057 row: 10
Class: transposon-tagged.
Location/Qualifiers
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/clone lib="1118 - RescueMu Grid S"
/note="Organ: leaf; Vector: RescueMu (engineered from
pBlueScript backbone); Site: 1: BamHI; Site 2: BglII;
RescueMu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
site 'www.zmdb.iastate.edu' and follow the links for
'RescueMu,' Grid S was grown at San Diego in 2002. DNA was
extracted from leaf strips, double digested using BamHI
and BglII, and ligated to form circular plasmids. DH10B
cells were transformed and then screened on LB plates with
ampicillin."

FEATURES
source

ORIGIN

Query Match 71.1%; Score 19.2; DB 9; Length 115;
Best Local Similarity 87.5%; Pred. No. 4.3e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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Db 58 GCCTGGCTACCCCTAGCCTAGGC 81
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RESULT 7

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DEFINITION
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survey sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Zea mays

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Walbot, V.
Unpublished (2001)
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site so sequence was trimmed. Post-ligation
sequence submitted separately.
Plate: 1118057 row: 10
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/clone lib="1118 - RescueMu Grid S"
/note="Organ: leaf; Vector: RescueMu (engineered from
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RescueMu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
site 'www.zmdb.iastate.edu' and follow the links for
'RescueMu,' Grid S was grown at San Diego in 2002. DNA was
extracted from leaf strips, double digested using BamHI
and BglII, and ligated to form circular plasmids. DH10B
cells were transformed and then screened on LB plates with
ampicillin."

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Walbot, V.
Unpublished (2001)
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site so sequence was trimmed. Post-ligation
sequence submitted separately.
Plate: 1118057 row: 10
Class: transposon-tagged.
Location/Qualifiers
1..115
/organism="Zea mays"
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/clone lib="1118 - RescueMu Grid S"
/note="Organ: leaf; Vector: RescueMu (engineered from
pBlueScript backbone); Site: 1: BamHI; Site 2: BglII;
RescueMu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
site 'www.zmdb.iastate.edu' and follow the links for
'RescueMu,' Grid S was grown at San Diego in 2002. DNA was
extracted from leaf strips, double digested using BamHI
and BglII, and ligated to form circular plasmids. DH10B
cells were transformed and then screened on LB plates with
ampicillin."

COMMENT

Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2297
Fax: 650 725 8221
Email: walbot@stanford.edu

Possible ligation site so sequence was trimmed. Post-ligation
sequence submitted separately.

Plate: 1118058 row: 10

Class: transposon-tagged

FEATURES

source

Location/Qualifiers

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/lab_host="DH10B"

/clone_lib="1118 -- RescueMu Grid S"

/note="Organ: leaf; Vector: RescueMu (engineered from

pBlueScript backbone); Site 1: BamHI; Site 2: BglII;

RescueMu is a 4.9 kb, modified maize Mu transposon

designed to allow plasmid rescue from total genomic DNA.

Mu elements insert preferentially into transcription

units. For more information on RescueMu, go to the web

site 'www.zmdb.iastate.edu' and follow the links for

'RescueMu.' Grid S was grown at San Diego in 2002. DNA was

extracted from leaf strips, double digested using BamHI

and BglII, and ligated to form circular plasmids. DH10B

cells were transformed and then screened on LB plates with

ampicillin."

ORIGIN

Query Match 71.1%; Score 19.2; DB 9; Length 115;

Best Local Similarity 87.5%; Pred. No. 4.3e+02;

Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 GCCTAGCTACCCCTAGGCTAGGC 27

DB 58 GCCTGGCTACCCCTAGGCTAGGC 81

RESULT 8

AZ636851

LOCUS

DEFINITION IM0495122R Mouse 10kb plasmid UUGC1M library Mus musculus genomic

clone UUGC1M0495122 R, genomic survey sequence.

AZ636851

VERSION AZ636851.1 GI:11759041

KEYWORDS GSS.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 419)

AUTHORS Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,

Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,

Kelly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von

Niederhauser, A. and Wright, D., Weiss, R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

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Insert Length: 10000 Std Error: 0.00

FEATURES

source

Location/Qualifiers

1..419

/organism="Mus musculus"

/mol_type="genomic DNA"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC1M0495122"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"

/clone_lib="Mouse 10kb plasmid UUGC1M library"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of pMD42 (GI|4732114|gb|AF129072.1), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

adapted vector DNA, and transformed into

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

ORIGIN

Query Match 71.1%; Score 19.2; DB 8; Length 419;

Best Local Similarity 87.5%; Pred. No. 4.5e+02;

Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 AGCCTAGCTACCCCTAGGCTAGG 26

DB 51 AGTCTAGCTTCCCTAGGTCAGG 74

RESULT 9

CC684396

LOCUS

DEFINITION CC684396 453 bp DNA linear GSS 19-JUN-2003

OSUKC11TV ZM 0.7_1.5_KB Zea mays genomic clone ZMWBMa0459B21,

genomic survey sequence.

CC684396

VERSION CC684396.1 GI:32089172

KEYWORDS GSS.

SOURCE Zea mays

ORGANISM Zea mays

REFERENCE 1 (bases 1 to 453)

AUTHORS Whitlaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,

Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,

Citek, R.W., Nurnberg, A., Robbins, D. and Lakey, N.

Consortium for Maize Genomics

Other GSSs: OSUKC11TH

Unpublished (2002)

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Seq primer: TP

Class: sheared ends.

Location/Qualifiers

FEATURES


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source
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/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone_lib="ZMMBta0459B21"
/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN
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Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCTA 24
|||||
Db 83 GTAGCCTAGTACCCCTAGCACCA 106

RESULT 10
BZ631002/c
LOCUS
DEFINITION
BZ631002 630 bp DNA linear GSS 29-JAN-2003
PAAZ280TD ZM_0.6_1.0_KB Zea mays genomic clone ZMMBta007M16,
genomic survey sequence.
ACCESSION
BZ631002
VERSION
BZ631002.1 GI:28077996
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 630)
Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Yuan,Y., San Miguel,P., Ma,J. and
Bennetzen,J.
Maize Genomics Consortium
Unpublished (2003)
Other GSSs: PAAZ280TB
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Email: whitelaw@tigr.org
Seq primer: TF
Class: sheared ends.
Location/Qualifiers
1. .672
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone_lib="ZMMBta007M16"
/clone_lib="ZM_0.6_1.0_KB"
/notes="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high
Cot selected genomic DNA library"

ORIGIN
Query Match 71.1%; Score 19.2; DB 8; Length 672;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCTA 24
|||||
Db 116 GTAGCCTATCCACCCCTAGGTGTA 139

RESULT 12
BZ633850
LOCUS
DEFINITION
BZ633850 683 bp DNA linear GSS 29-JAN-2003
PUAY40TB ZM_0.6_1.0_KB Zea mays genomic clone ZMMBta007G07,
genomic survey sequence.
ACCESSION
BZ633850
VERSION
BZ633850.1 GI:28080844
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 683)
Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Yuan,Y., San Miguel,P., Ma,J. and
Bennetzen,J.
Maize Genomics Consortium
Unpublished (2003)
Other GSSs: PUAY40TD
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Email: whitelaw@tigr.org
Seq primer: TF
Class: sheared ends.

ORIGIN
Query Match 71.1%; Score 19.2; DB 8; Length 630;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GTAGCCTAGTACCCCTAGGTCTA 24
|||||
Db 472 GTAGCCTATCCACCCCTAGGTGTA 449

RESULT 11
BZ630997
LOCUS
DEFINITION
BZ630997 672 bp DNA linear GSS 29-JAN-2003
PAAZ280TB ZM_0.6_1.0_KB Zea mays genomic clone ZMMBta007M16,
genomic survey sequence.
ACCESSION
BZ630997
VERSION
BZ630997.1 GI:28077991
KEYWORDS
GSS.
SOURCE
Zea mays
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 672)
Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Yuan,Y., San Miguel,P., Ma,J. and
Bennetzen,J.
Maize Genomics Consortium
Unpublished (2003)
Other GSSs: PAAZ280TD
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Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TF
Class: sheared ends.

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FEATURES
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      /note="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high
      Cot selected genomic DNA library"

ORIGIN
Query Match      71.1%; Score 19.2; DB 8; Length 683;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GTAGCCTAGCTACCCCTAGGTCTA 24
    |||||
Db 116 GTAGCCTATCCACCCTAGGTGTA 139

RESULT 13
LOCUS BH837552/c 705 bp DNA linear GSS 28-MAY-2002
DEFINITION LMCRI00001A03f Zea mays L. Zea mays genomic clone LMCRI00001A03f,
genomic survey sequence.
ACCESSION BH837552
VERSION BH837552.1 GI:21235430
KEYWORDS GSS.
SOURCE Zea mays
  ORGANISM Zea mays
    Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
    Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
    clade; Panicoideae; Andropogoneae; Zea.
  1 (bases 1 to 705)
  Kim, S.W., Yu, Y., Lee, M.C., Main, D. and Wing, R.A.
  Methyl-filtration genomic sequence from maize
  Unpublished (2002)
  Contact: Wing RA
  Clemson University Genomics Institute
  100 Jordan Hall, Clemson, SC 29634, USA
  Tel: 864 656 7288
  Fax: 864 656 4293
  Email: rwing@clemson.edu
  Total High Quality bases = 551
  Seq primer: TAATACGACTCACTATAGGG
  Class: shotgun
  High quality sequence start: 12
  High quality sequence stop: 690.

FEATURES
  source
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      /tissue_type="Leaf"
      /lab_host="DH10B"
      /clone_lib="Zea mays L."
      /note="Vector: pGEM-T easy; Site 1: Mcr BC;
      Methyl-filtration library, Nuclei DNA was completely
      digested with Mcr BC, size fractionated and transformed
      to E.Coli.DH10B."

ORIGIN
Query Match      71.1%; Score 19.2; DB 8; Length 705;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 4 GCCTAGCTACCCCTAGGTCTAGGC 27
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Db 128 GCCTGGCTACCCCTAGCCCTAGGC 105

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RESULT 14
LOCUS BZ633854/c 725 bp DNA linear GSS 29-JAN-2003
DEFINITION PUAY40TD ZM 0.6-1.0 KB Zea mays genomic clone ZMMBTA007G07,
genomic survey sequence.
ACCESSION BZ633854
VERSION BZ633854.1 GI:28080848
KEYWORDS GSS.
SOURCE Zea mays
  ORGANISM Zea mays
    Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
    Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
    clade; Panicoideae; Andropogoneae; Zea.
  1 (bases 1 to 725)
  White, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
  Resnick, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and
  Bennetzen, J.
  Maize Genomics Consortium
  Unpublished (2003)
  Other_GSSs: PUAY40TB
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  TIGR
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  Email: whitelaw@tigr.org
  Seq primer: TF
  Class: sheared ends.

FEATURES
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      /mol_type="genomic DNA"
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      /db_xref="taxon:4577"
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      Cot selected genomic DNA library"

ORIGIN
Query Match      71.1%; Score 19.2; DB 8; Length 725;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GTAGCCTAGCTACCCCTAGGTCTA 24
    |||||
Db 567 GTAGCCTATCCACCCTAGGTGTA 544

RESULT 15
LOCUS CG351990/c 745 bp DNA linear GSS 26-AUG-2003
DEFINITION OGVC84TH ZM 0.7-1.5 KB Zea mays genomic clone ZMMBWA0504N23,
genomic survey sequence.
ACCESSION CG351990
VERSION CG351990.1 GI:34269256
KEYWORDS GSS.
SOURCE Zea mays
  ORGANISM Zea mays
    Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
    Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
    clade; Panicoideae; Andropogoneae; Zea.
  1 (bases 1 to 745)
  White, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
  Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,
  Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.
  Consortium for Maize Genomics
  Unpublished (2002)
  Other_GSSs: OGVC84TV
  Contact: Cathy Whitelaw
  TIGR

```

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Seq primer: TR

Class: Sheared ends.

FEATURES

source

Location/Qualifiers

1. .745

/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

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/clone_lib="ZM_0.7-1.5_KB"

/notes="Vector: pBCSK-; Site_1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN

Query Match 71.1%; Score 19.2; DB 9; Length 745;
Best Local Similarity 87.5%; Pred. No. 4.5e+02;
Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 GCCTAGCTACCCCTAGCTAGGC 27

Db 389 GCCTGGCTACCCCTAGCCCTAGGC 366

Search completed: October 13, 2005, 21:43:33

Job time : 7564 secs